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(54) Title: NOVEL NUCLEIC ACIDS AND POLYPEPTIDES

(57) Abstract: The present invention provides nucleic acids, novel polypeptide sequences encoded by these nucleic acids and uses thereof.

NOVEL NUCLEIC ACIDS AND POLYPEPTIDES

1. TECHNICAL FIELD

The present invention provides novel polynucleotides and proteins encoded by such polynucleotides, along with uses for these polynucleotides and proteins, for example in therapeutic, diagnostic and research methods.

2. BACKGROUND

Technology aimed at the discovery of protein factors (including e.g., cytokines, such as lymphokines, interferons, circulating soluble factors, chemokines, and interleukins) has matured rapidly over the past decade. The now routine hybridization cloning and expression cloning techniques clone novel polynucleotides "directly" in the sense that they rely on information directly related to the discovered protein (i.e., partial DNA/amino acid sequence of the protein in the case of hybridization cloning; activity of the protein in the case of expression cloning). More recent "indirect" cloning techniques such as signal sequence cloning, which isolates DNA sequences based on the presence of a now well-recognized secretory leader sequence motif, as well as various PCR-based or low stringency hybridization-based cloning techniques, have advanced the state of the art by making available large numbers of DNA/amino acid sequences for proteins that are known to have biological activity, for example, by virtue of their secreted nature in the case of leader sequence cloning, by virtue of their cell or tissue source in the case of PCR-based techniques, or by virtue of structural similarity to other genes of known biological activity.

Identified polynucleotide and polypeptide sequences have numerous applications in, for example, diagnostics, forensics, gene mapping; identification of mutations responsible for genetic disorders or other traits, to assess biodiversity, and to produce many other types of data and products dependent on DNA and amino acid sequences.

3. SUMMARY OF THE INVENTION

The compositions of the present invention include novel isolated polypeptides, novel isolated polypucleotides encoding such polypeptides, including recombinant DNA molecules, cloned genes or degenerate variants thereof, especially naturally occurring variants such as allelic variants, antisense polynucleotide molecules, and antibodies that specifically recognize one or more epitopes present on such polypeptides, as well as hybridomas producing such antibodies.

The compositions of the present invention additionally include vectors, including expression vectors, containing the polynucleotides of the invention, cells genetically engineered to contain such polynucleotides and cells genetically engineered to express such polynucleotides.

The present invention relates to a collection or library of at least one novel nucleic acid sequence assembled from expressed sequence tags (ESTs) isolated mainly by sequencing by hybridization (SBH), and in some cases, sequences obtained from one or more public databases. The invention relates also to the proteins encoded by such polynucleotides, along with therapeutic, diagnostic and research utilities for these polynucleotides and proteins. These nucleic acid sequences are designated as SEQ ID NO: 1-444. The polypeptides sequences are designated SEQ ID NO: 445-888. The nucleic acids and polypeptides are provided in the Sequence Listing. In the nucleic acids provided in the Sequence Listing, A is adenosine; C is cytosine; G is guanine; T is thymine; and N is unknown or any of the four bases.

The nucleic acid sequences of the present invention also include, nucleic acid sequences that hybridize to the complement of SEQ ID NO: 1-444 under stringent hybridization conditions; nucleic acid sequences which are allelic variants or species homologues of any of the nucleic acid sequences recited above, or nucleic acid sequences that encode a peptide comprising a specific domain or truncation of the peptides encoded by SEQ ID NO: 1-444. A polynucleotide comprising a nucleotide sequence having at least 90% identity to an identifying sequence of SEQ ID NO: 1-444 or a degenerate variant or fragment thereof. The identifying sequence can be 100 base pairs in length.

The nucleic acid sequences of the present invention also include the sequence information from the nucleic acid sequences of SEQ ID NO: 1-444. The sequence information can be a segment of any one of SEQ ID NO: 1-444 that uniquely identifies or represents the sequence information of SEQ ID NO: 1-444.

A collection as used in this application can be a collection of only one polynucleotide. The collection of sequence information or identifying information of each sequence can be provided on a nucleic acid array. In one embodiment, segments of sequence information are provided on a nucleic acid array to detect the polynucleotide that contains the segment. The array can be designed to detect full-match or mismatch to the polynucleotide that contains the segment. The collection can also be provided in a computer-readable format.

This invention also includes the reverse or direct complement of any of the nucleic acid sequences recited above; cloning or expression vectors containing the nucleic acid sequences; and host cells or organisms transformed with these expression vectors. Nucleic acid sequences (or their reverse or direct complements) according to the invention have numerous applications in a variety of techniques known to those skilled in the art of molecular biology, such as use as hybridization

probes, use as primers for PCR, use in an array, use in computer-readable media, use in sequencing full-length genes, use for chromosome and gene mapping, use in the recombinant production of protein, and use in the generation of anti-sense DNA or RNA, their chemical analogs and the like.

In a preferred embodiment, the nucleic acid sequences of SEQ ID NO: 1-444 or novel segments or parts of the nucleic acids of the invention are used as primers in expression assays that are well known in the art. In a particularly preferred embodiment, the nucleic acid sequences of SEQ ID NO: 1-444 or novel segments or parts of the nucleic acids provided herein are used in diagnostics for identifying expressed genes or, as well known in the art and exemplified by Vollrath et al., Science 258:52-59 (1992), as expressed sequence tags for physical mapping of the human genome.

The isolated polynucleotides of the invention include, but are not limited to, a polynucleotide comprising any one of the nucleotide sequences set forth in SEQ ID NO: 1-444; a polynucleotide comprising any of the full length protein coding sequences of SEQ ID NO: 1-444; and a polynucleotide comprising any of the nucleotide sequences of the mature protein coding sequences of SEQ ID NO: 1-444. The polynucleotides of the present invention also include, but are not limited to, a polynucleotide that hybridizes under stringent hybridization conditions to (a) the complement of any one of the nucleotide sequences set forth in SEQ ID NO: 1-444; (b) a nucleotide sequence encoding any one of the amino acid sequences set forth in the Sequence Listing; (c) a polynucleotide which is an allelic variant of any polynucleotides recited above; (d) a polynucleotide which encodes a species homolog (e.g. orthologs) of any of the proteins recited above; or (e) a polynucleotide that encodes a polypeptide comprising a specific domain or truncation of any of the polypeptides comprising an amino acid sequence set forth in the Sequence Listing.

The isolated polypeptides of the invention include, but are not limited to, a polypeptide comprising any of the amino acid sequences set forth in SEQ ID NO: 445-888; or the corresponding full length or mature protein. Polypeptides of the invention also include polypeptides with biological activity that are encoded by (a) any of the polynucleotides having a nucleotide sequence set forth in SEQ ID NO: 1-444; or (b) polynucleotides that hybridize to the complement of the polynucleotides of (a) under stringent hybridization conditions. Biologically or immunologically active variants of any of the polypeptide sequences in the Sequence Listing, and "substantial equivalents" thereof (e.g., with at least about 65%, 70%, 75%, 80%, 85%, 90%, 95%, 98% or 99% amino acid sequence identity) that preferably retain biological activity are also contemplated. The polypeptides of the invention may be wholly or partially chemically synthesized but are preferably produced by recombinant means using the genetically engineered cells (e.g. host cells) of the invention.

hydrophilic, e.g., pharmaceutically acceptable, carrier.

The invention also provides compositions comprising a polypeptide of the invention.

Polypeptide compositions of the invention may further comprise an acceptable carrier, such as a

The invention also provides host cells transformed or transfected with a polynucleotide of the invention.

The invention also relates to methods for producing a polypeptide of the invention comprising growing a culture of the host cells of the invention in a suitable culture medium under conditions permitting expression of the desired polypeptide, and purifying the polypeptide from the culture or from the host cells. Preferred embodiments include those in which the protein produced by such process is a mature form of the protein.

Polynucleotides according to the invention have numerous applications in a variety of techniques known to those skilled in the art of molecular biology. These techniques include use as hybridization probes, use as oligomers, or primers, for PCR, use for chromosome and gene mapping, use in the recombinant production of protein, and use in generation of anti-sense DNA or RNA, their chemical analogs and the like. For example, when the expression of an mRNA is largely restricted to a particular cell or tissue type, polynucleotides of the invention can be used as hybridization probes to detect the presence of the particular cell or tissue mRNA in a sample using, e.g., in situ hybridization.

In other exemplary embodiments, the polynucleotides are used in diagnostics as expressed sequence tags for identifying expressed genes or, as well known in the art and exemplified by Vollrath et al., Science 258:52-59 (1992), as expressed sequence tags for physical mapping of the human genome.

The polypeptides according to the invention can be used in a variety of conventional procedures and methods that are currently applied to other proteins. For example, a polypeptide of the invention can be used to generate an antibody that specifically binds the polypeptide. Such antibodies, particularly monoclonal antibodies, are useful for detecting or quantitating the polypeptide in tissue. The polypeptides of the invention can also be used as molecular weight markers, and as a food supplement.

Methods are also provided for preventing, treating, or ameliorating a medical condition which comprises the step of administering to a mammalian subject a therapeutically effective amount of a composition comprising a polypeptide of the present invention and a pharmaceutically acceptable carrier.

In particular, the polypeptides and polynucleotides of the invention can be utilized, for example, in methods for the prevention and/or treatment of disorders involving aberrant protein expression or biological activity.

The present invention further relates to methods for detecting the presence of the polynucleotides or polypeptides of the invention in a sample. Such methods can, for example, be utilized as part of prognostic and diagnostic evaluation of disorders as recited herein and for the identification of subjects exhibiting a predisposition to such conditions. The invention provides a method for detecting the polynucleotides of the invention in a sample, comprising contacting the sample with a compound that binds to and forms a complex with the polynucleotide of interest for a period sufficient to form the complex and under conditions sufficient to form a complex and detecting the complex such that if a complex is detected, the polynucleotide of interest is detected. The invention also provides a method for detecting the polypeptides of the invention in a sample comprising contacting the sample with a compound that binds to and forms a complex with the polypeptide under conditions and for a period sufficient to form the complex and detecting the formation of the complex such that if a complex is formed, the polypeptide is detected.

The invention also provides kits comprising polynucleotide probes and/or monoclonal antibodies, and optionally quantitative standards, for carrying out methods of the invention. Furthermore, the invention provides methods for evaluating the efficacy of drugs, and monitoring the progress of patients, involved in clinical trials for the treatment of disorders as recited above.

The invention also provides methods for the identification of compounds that modulate (i.e., increase or decrease) the expression or activity of the polynucleotides and/or polypeptides of the invention. Such methods can be utilized, for example, for the identification of compounds that can ameliorate symptoms of disorders as recited herein. Such methods can include, but are not limited to, assays for identifying compounds and other substances that interact with (e.g., bind to) the polypeptides of the invention. The invention provides a method for identifying a compound that binds to the polypeptides of the invention comprising contacting the compound with a polypeptide of the invention in a cell for a time sufficient to form a polypeptide/compound complex, wherein the complex drives expression of a reporter gene sequence in the cell; and detecting the complex by detecting the reporter gene sequence expression such that if expression of the reporter gene is detected the compound the binds to a polypeptide of the invention is identified.

The methods of the invention also provide methods for treatment which involve the administration of the polynucleotides or polypeptides of the invention to individuals exhibiting symptoms or tendencies. In addition, the invention encompasses methods for treating diseases or disorders as recited herein comprising administering compounds and other substances that modulate the overall activity of the target gene products. Compounds and other substances can

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effect such modulation either on the level of target gene/protein expression or target protein activity.

The polypeptides of the present invention and the polynucleotides encoding them are also useful for the same functions known to one of skill in the art as the polypeptides and polynucleotides to which they have homology (set forth in Table 2); for which they have a signature region (as set forth in Table 3); or for which they have homology to a gene family (as set forth in Table 4). If no homology is set forth for a sequence, then the polypeptides and polynucleotides of the present invention are useful for a variety of applications, as described herein, including use in arrays for detection.

4. DETAILED DESCRIPTION OF THE INVENTION

4.1 DEFINITIONS

It must be noted that as used herein and in the appended claims, the singular forms "a", "an" and "the" include plural references unless the context clearly dictates otherwise.

The term "active" refers to those forms of the polypeptide which retain the biologic and/or immunologic activities of any naturally occurring polypeptide. According to the invention, the terms "biologically active" or "biological activity" refer to a protein or peptide having structural, regulatory or biochemical functions of a naturally occurring molecule. Likewise "immunologically active" or "immunological activity" refers to the capability of the natural, recombinant or synthetic polypeptide to induce a specific immune response in appropriate animals or cells and to bind with specific antibodies.

The term "activated cells" as used in this application are those cells which are engaged in extracellular or intracellular membrane trafficking, including the export of secretory or enzymatic molecules as part of a normal or disease process.

The terms "complementary" or "complementarity" refer to the natural binding of polynucleotides by base pairing. For example, the sequence 5'-AGT-3' binds to the complementary sequence 3'-TCA-5'. Complementarity between two single-stranded molecules may be "partial" such that only some of the nucleic acids bind or it may be "complete" such that total complementarity exists between the single stranded molecules. The degree of complementarity between the nucleic acid strands has significant effects on the efficiency and strength of the hybridization between the nucleic acid strands.

The term "embryonic stem cells (ES)" refers to a cell that can give rise to many differentiated cell types in an embryo or an adult, including the germ cells. The term "germ line stem cells (GSCs)" refers to stem cells derived from primordial stem cells that provide a steady

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and continuous source of germ cells for the production of gametes. The term "primordial germ cells (PGCs)" refers to a small population of cells set aside from other cell lineages particularly from the yolk sac, mesenteries, or gonadal ridges during embryogenesis that have the potential to differentiate into germ cells and other cells. PGCs are the source from which GSCs and ES cells are derived. The PGCs, the GSCs and the ES cells are capable of self-renewal. Thus these cells not only populate the germ line and give rise to a plurality of terminally differentiated cells that comprise the adult specialized organs, but are able to regenerate themselves.

The term "expression modulating fragment," EMF, means a series of nucleotides which modulates the expression of an operably linked ORF or another EMF.

As used herein, a sequence is said to "modulate the expression of an operably linked sequence" when the expression of the sequence is altered by the presence of the EMF. EMFs include, but are not limited to, promoters, and promoter modulating sequences (inducible elements). One class of EMFs are nucleic acid fragments which induce the expression of an operably linked ORF in response to a specific regulatory factor or physiological event.

The terms "nucleotide sequence" or "nucleic acid" or "polynucleotide" or "oligonculeotide" are used interchangeably and refer to a heteropolymer of nucleotides or the sequence of these nucleotides. These phrases also refer to DNA or RNA of genomic or synthetic origin which may be single-stranded or double-stranded and may represent the sense or the antisense strand, to peptide nucleic acid (PNA) or to any DNA-like or RNA-like material. In the sequences herein A is adenine, C is cytosine, T is thymine, G is guanine and N is A, C, G or T (U). It is contemplated that where the polynucleotide is RNA, the T (thymine) in the sequences provided herein is substituted with U (uracil). Generally, nucleic acid segments provided by this invention may be assembled from fragments of the genome and short oligonucleotide linkers, or from a series of oligonucleotides, or from individual nucleotides, to provide a synthetic nucleic acid which is capable of being expressed in a recombinant transcriptional unit comprising regulatory elements derived from a microbial or viral operon, or a eukaryotic gene.

The terms "oligonucleotide fragment" or a "polynucleotide fragment", "portion," or "segment" or "probe" or "primer" are used interchangeably and refer to a sequence of nucleotide residues which are at least about 5 nucleotides, more preferably at least about 7 nucleotides, more preferably at least about 9 nucleotides, more preferably at least about 11 nucleotides and most preferably at least about 17 nucleotides. The fragment is preferably less than about 500 nucleotides, preferably less than about 200 nucleotides, more preferably less than about 100 nucleotides, more preferably less than about 50 nucleotides and most preferably less than 30 nucleotides. Preferably the probe is from about 6 nucleotides to about 200 nucleotides, preferably from about 15 to about 50 nucleotides, more preferably from about 17 to 30

nucleotides and most preferably from about 20 to 25 nucleotides. Preferably the fragments can be used in polymerase chain reaction (PCR), various hybridization procedures or microarray procedures to identify or amplify identical or related parts of mRNA or DNA molecules. A fragment or segment may uniquely identify each polynucleotide sequence of the present invention. Preferably the fragment comprises a sequence substantially similar to any one of SEQ ID NO: 1-444.

Probes may, for example, be used to determine whether specific mRNA molecules are present in a cell or tissue or to isolate similar nucleic acid sequences from chromosomal DNA as described by Walsh et al. (Walsh, P.S. et al., 1992, PCR Methods Appl 1:241-250). They may be labeled by nick translation, Klenow fill-in reaction, PCR, or other methods well known in the art. Probes of the present invention, their preparation and/or labeling are elaborated in Sambrook, J. et al., 1989, Molecular Cloning: A Laboratory Manual, Cold Spring Harbor Laboratory, NY; or Ausubel, F.M. et al., 1989, Current Protocols in Molecular Biology, John Wiley & Sons, New York NY, both of which are incorporated herein by reference in their entirety.

The nucleic acid sequences of the present invention also include the sequence information from the nucleic acid sequences of SEQ ID NO: 1-444. The sequence information can be a segment of any one of SEQ ID NO: 1-444 that uniquely identifies or represents the sequence information of that sequence of SEQ ID NO: 1-444. One such segment can be a twenty-mer nucleic acid sequence because the probability that a twenty-mer is fully matched in the human genome is 1 in 300. In the human genome, there are three billion base pairs in one set of chromosomes. Because 4²⁰ possible twenty-mers exist, there are 300 times more twenty-mers than there are base pairs in a set of human chromosomes. Using the same analysis, the probability for a seventeen-mer to be fully matched in the human genome is approximately 1 in 5. When these segments are used in arrays for expression studies, fifteen-mer segments can be used. The probability that the fifteen-mer is fully matched in the expressed sequences is also approximately one in five because expressed sequences comprise less than approximately 5% of the entire genome sequence.

Similarly, when using sequence information for detecting a single mismatch, a segment can be a twenty-five mer. The probability that the twenty-five mer would appear in a human genome with a single mismatch is calculated by multiplying the probability for a full match $(1 \div 4^{25})$ times the increased probability for mismatch at each nucleotide position (3×25) . The probability that an eighteen mer with a single mismatch can be detected in an array for expression studies is approximately one in five. The probability that a twenty-mer with a single mismatch can be detected in a human genome is approximately one in five.

The term "open reading frame," ORF, means a series of nucleotide triplets coding for amino acids without any termination codons and is a sequence translatable into protein.

The terms "operably linked" or "operably associated" refer to functionally related nucleic acid sequences. For example, a promoter is operably associated or operably linked with a coding sequence if the promoter controls the transcription of the coding sequence. While operably linked nucleic acid sequences can be contiguous and in the same reading frame, certain genetic elements e.g. repressor genes are not contiguously linked to the coding sequence but still control transcription/translation of the coding sequence.

The term "pluripotent" refers to the capability of a cell to differentiate into a number of differentiated cell types that are present in an adult organism. A pluripotent cell is restricted in its differentiation capability in comparison to a totipotent cell.

The terms "polypeptide" or "peptide" or "amino acid sequence" refer to an oligopeptide, peptide, polypeptide or protein sequence or fragment thereof and to naturally occurring or synthetic molecules. A polypeptide "fragment," "portion," or "segment" is a stretch of amino acid residues of at least about 5 amino acids, preferably at least about 7 amino acids, more preferably at least about 9 amino acids and most preferably at least about 17 or more amino acids. The peptide preferably is not greater than about 500 amino acids, more preferably less than 200 amino acids more preferably less than 150 amino acids and most preferably less than 100 amino acids. Preferably the peptide is from about 5 to about 200 amino acids. To be active, any polypeptide must have sufficient length to display biological and/or immunological activity.

The term "naturally occurring polypeptide" refers to polypeptides produced by cells that have not been genetically engineered and specifically contemplates various polypeptides arising from post-translational modifications of the polypeptide including, but not limited to, acetylation, carboxylation, glycosylation, phosphorylation, lipidation and acylation.

The term "translated protein coding portion" means a sequence which encodes for the full length protein which may include any leader sequence or any processing sequence.

The term "mature protein coding sequence" means a sequence which encodes a peptide or protein without a signal or leader sequence. The "mature protein portion" means that portion of the protein which does not include a signal or leader sequence. The peptide may have been produced by processing in the cell which removes any leader/signal sequence. The mature protein portion may or may not include an initial methionine residue. The methionine residue may be removed from the protein during processing in the cell. The peptide may be produced synthetically or the protein may have been produced using a polynucleotide only encoding for the mature protein coding sequence.

The term "derivative" refers to polypeptides chemically modified by such techniques as ubiquitination, labeling (e.g., with radionuclides or various enzymes), covalent polymer attachment such as pegylation (derivatization with polyethylene glycol) and insertion or substitution by chemical synthesis of amino acids such as ornithine, which do not normally occur in human proteins.

The term "variant" (or "analog") refers to any polypeptide differing from naturally occurring polypeptides by amino acid insertions, deletions, and substitutions, created using, e.g., recombinant DNA techniques. Guidance in determining which amino acid residues may be replaced, added or deleted without abolishing activities of interest, may be found by comparing the sequence of the particular polypeptide with that of homologous peptides and minimizing the number of amino acid sequence changes made in regions of high homology (conserved regions) or by replacing amino acids with consensus sequence.

Alternatively, recombinant variants encoding these same or similar polypeptides may be synthesized or selected by making use of the "redundancy" in the genetic code. Various codon substitutions, such as the silent changes which produce various restriction sites, may be introduced to optimize cloning into a plasmid or viral vector or expression in a particular prokaryotic or eukaryotic system. Mutations in the polynucleotide sequence may be reflected in the polypeptide or domains of other peptides added to the polypeptide to modify the properties of any part of the polypeptide, to change characteristics such as ligand-binding affinities, interchain affinities, or degradation/turnover rate.

Preferably, amino acid "substitutions" are the result of replacing one amino acid with another amino acid having similar structural and/or chemical properties, i.e., conservative amino acid replacements. "Conservative" amino acid substitutions may be made on the basis of similarity in polarity, charge, solubility, hydrophobicity, hydrophilicity, and/or the amphipathic nature of the residues involved. For example, nonpolar (hydrophobic) amino acids include alanine, leucine, isoleucine, valine, proline, phenylalanine, tryptophan, and methionine; polar neutral amino acids include glycine, serine, threonine, cysteine, tyrosine, asparagine, and glutamine; positively charged (basic) amino acids include arginine, lysine, and histidine; and negatively charged (acidic) amino acids include aspartic acid and glutamic acid. "Insertions" or "deletions" are preferably in the range of about 1 to 20 amino acids, more preferably 1 to 10 amino acids. The variation allowed may be experimentally determined by systematically making insertions, deletions, or substitutions of amino acids in a polypeptide molecule using recombinant DNA techniques and assaying the resulting recombinant variants for activity.

Alternatively, where alteration of function is desired, insertions, deletions or non-conservative alterations can be engineered to produce altered polypeptides. Such alterations

can, for example, alter one or more of the biological functions or biochemical characteristics of the polypeptides of the invention. For example, such alterations may change polypeptide characteristics such as ligand-binding affinities, interchain affinities, or degradation/turnover rate. Further, such alterations can be selected so as to generate polypeptides that are better suited for expression, scale up and the like in the host cells chosen for expression. For example, cysteine residues can be deleted or substituted with another amino acid residue in order to eliminate disulfide bridges.

The terms "purified" or "substantially purified" as used herein denotes that the indicated nucleic acid or polypeptide is present in the substantial absence of other biological macromolecules, e.g., polynucleotides, proteins, and the like. In one embodiment, the polynucleotide or polypeptide is purified such that it constitutes at least 95% by weight, more preferably at least 99% by weight, of the indicated biological macromolecules present (but water, buffers, and other small molecules, especially molecules having a molecular weight of less than 1000 daltons, can be present).

The term "isolated" as used herein refers to a nucleic acid or polypeptide separated from at least one other component (e.g., nucleic acid or polypeptide) present with the nucleic acid or polypeptide in its natural source. In one embodiment, the nucleic acid or polypeptide is found in the presence of (if anything) only a solvent, buffer, ion, or other component normally present in a solution of the same. The terms "isolated" and "purified" do not encompass nucleic acids or polypeptides present in their natural source.

The term "recombinant," when used herein to refer to a polypeptide or protein, means that a polypeptide or protein is derived from recombinant (e.g., microbial, insect, or mammalian) expression systems. "Microbial" refers to recombinant polypeptides or proteins made in bacterial or fungal (e.g., yeast) expression systems. As a product, "recombinant microbial" defines a polypeptide or protein essentially free of native endogenous substances and unaccompanied by associated native glycosylation. Polypeptides or proteins expressed in most bacterial cultures, e.g., E. coli, will be free of glycosylation modifications; polypeptides or proteins expressed in yeast will have a glycosylation pattern in general different from those expressed in mammalian cells.

The term "recombinant expression vehicle or vector" refers to a plasmid or phage or virus or vector, for expressing a polypeptide from a DNA (RNA) sequence. An expression vehicle can comprise a transcriptional unit comprising an assembly of (1) a genetic element or elements having a regulatory role in gene expression, for example, promoters or enhancers, (2) a structural or coding sequence which is transcribed into mRNA and translated into protein, and (3) appropriate transcription initiation and termination sequences. Structural units intended for use

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in yeast or eukaryotic expression systems preferably include a leader sequence enabling extracellular secretion of translated protein by a host cell. Alternatively, where recombinant protein is expressed without a leader or transport sequence, it may include an amino terminal methionine residue. This residue may or may not be subsequently cleaved from the expressed recombinant protein to provide a final product.

The term "recombinant expression system" means host cells which have stably integrated a recombinant transcriptional unit into chromosomal DNA or carry the recombinant transcriptional unit extrachromosomally. Recombinant expression systems as defined herein will express heterologous polypeptides or proteins upon induction of the regulatory elements linked to the DNA segment or synthetic gene to be expressed. This term also means host cells which have stably integrated a recombinant genetic element or elements having a regulatory role in gene expression, for example, promoters or enhancers. Recombinant expression systems as defined herein will express polypeptides or proteins endogenous to the cell upon induction of the regulatory elements linked to the endogenous DNA segment or gene to be expressed. The cells can be prokaryotic or eukaryotic.

The term "secreted" includes a protein that is transported across or through a membrane, including transport as a result of signal sequences in its amino acid sequence when it is expressed in a suitable host cell. "Secreted" proteins include without limitation proteins secreted wholly (e.g., soluble proteins) or partially (e.g., receptors) from the cell in which they are expressed. "Secreted" proteins also include without limitation proteins that are transported across the membrane of the endoplasmic reticulum. "Secreted" proteins are also intended to include proteins containing non-typical signal sequences (e.g. Interleukin-1 Beta, see Krasney, P.A. and Young, P.R. (1992) Cytokine 4(2): 134-143) and factors released from damaged cells (e.g. Interleukin-1 Receptor Antagonist, see Arend, W.P. et. al. (1998) Annu. Rev. Immunol. 16:27-55)

Where desired; an expression vector may be designed to contain a "signal or leader sequence" which will direct the polypeptide through the membrane of a cell. Such a sequence may be naturally present on the polypeptides of the present invention or provided from heterologous protein sources by recombinant DNA techniques.

The term "stringent" is used to refer to conditions that are commonly understood in the art as stringent. Stringent conditions can include highly stringent conditions (i.e., hybridization to filter-bound DNA in 0.5 M NaHPO4, 7% sodium dodecyl sulfate (SDS), 1 mM EDTA at 65°C, and washing in 0.1X SSC/0.1% SDS at 68°C), and moderately stringent conditions (i.e., washing in 0.2X SSC/0.1% SDS at 42°C). Other exemplary hybridization conditions are described herein in the examples.

In instances of hybridization of deoxyoligonucleotides, additional exemplary stringent hybridization conditions include washing in 6X SSC/0.05% sodium pyrophosphate at 37°C (for 14-base oligonucleotides), 48°C (for 17-base oligos), 55°C (for 20-base oligonucleotides), and 60°C (for 23-base oligonucleotides).

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As used herein, "substantially equivalent" can refer both to nucleotide and amino acid sequences, for example a mutant sequence, that varies from a reference sequence by one or more substitutions, deletions, or additions, the net effect of which does not result in an adverse functional dissimilarity between the reference and subject sequences. Typically, such a substantially equivalent sequence varies from one of those listed herein by no more than about 35% (i.e., the number of individual residue substitutions, additions, and/or deletions in a substantially equivalent sequence, as compared to the corresponding reference sequence, divided by the total number of residues in the substantially equivalent sequence is about 0.35 or less). Such a sequence is said to have 65% sequence identity to the listed sequence. In one embodiment, a substantially equivalent, e.g., mutant, sequence of the invention varies from a listed sequence by no more than 30% (70% sequence identity); in a variation of this embodiment, by no more than 25% (75% sequence identity); and in a further variation of this embodiment, by no more than 20% (80% sequence identity) and in a further variation of this embodiment, by no more than 10% (90% sequence identity) and in a further variation of this embodiment, by no more that 5% (95% sequence identity). Substantially equivalent, e.g., mutant, amino acid sequences according to the invention preferably have at least 80% sequence identity with a listed amino acid sequence, more preferably at least 85% sequence identity, more preferably at least 90% sequence identity, more preferably at least 95% identity, more preferably at least 98% identity, and most preferably at least 99% identity. Substantially equivalent nucleotide sequences of the invention can have lower percent sequence identities, taking into account, for example, the redundancy or degeneracy of the genetic code. Preferably, nucleotide sequence has at least about 65% identity, more preferably at least about 75% identity, more preferably at least about 80% sequence identity, more preferably at least about 85% sequence identity, more preferably at least about 90% sequence identity, and most preferably at least about 95% identity, more preferably at least about 98% sequence identity, and most preferably at least about 99% sequence identity. For the purposes of the present invention, sequences having substantially equivalent biological activity and substantially equivalent expression characteristics are considered substantially equivalent. For the purposes of determining equivalence, truncation of the mature sequence (e.g., via a mutation which creates a spurious stop codon) should be disregarded. Sequence identity may be determined, e.g., using the Jotun Hein method (Hein, J.

(1990) Methods Enzymol. 183:626-645). Identity between sequences can also be determined by other methods known in the art, e.g. by varying hybridization conditions.

The term "totipotent" refers to the capability of a cell to differentiate into all of the cell types of an adult organism.

The term "transformation" means introducing DNA into a suitable host cell so that the DNA is replicable, either as an extrachromosomal element, or by chromosomal integration. The term "transfection" refers to the taking up of an expression vector by a suitable host cell, whether or not any coding sequences are in fact expressed. The term "infection" refers to the introduction of nucleic acids into a suitable host cell by use of a virus or viral vector.

As used herein, an "uptake modulating fragment," UMF, means a series of nucleotides which mediate the uptake of a linked DNA fragment into a cell. UMFs can be readily identified using known UMFs as a target sequence or target motif with the computer-based systems described below. The presence and activity of a UMF can be confirmed by attaching the suspected UMF to a marker sequence. The resulting nucleic acid molecule is then incubated with an appropriate host under appropriate conditions and the uptake of the marker sequence is determined. As described above, a UMF will increase the frequency of uptake of a linked marker sequence.

Each of the above terms is meant to encompass all that is described for each, unless the context dictates otherwise.

4.2 NUCLEIC ACIDS OF THE INVENTION

Nucleotide sequences of the invention are set forth in the Sequence Listing.

The isolated polynucleotides of the invention include a polynucleotide comprising the nucleotide sequences of SEQ ID NO: 1-444; a polynucleotide encoding any one of the peptide sequences of SEQ ID NO: 445-888; and a polynucleotide comprising the nucleotide sequence encoding the mature protein coding sequence of the polypeptides of any one of SEQ ID NO: 445-888. The polynucleotides of the present invention also include, but are not limited to, a polynucleotide that hybridizes under stringent conditions to (a) the complement of any of the nucleotides sequences of SEQ ID NO: 1-444; (b) nucleotide sequences encoding any one of the amino acid sequences set forth in the Sequence Listing as SEQ ID NO: 445-888; (c) a polynucleotide which is an allelic variant of any polynucleotide recited above; (d) a polynucleotide which encodes a species homolog of any of the proteins recited above; or (e) a polynucleotide that encodes a polypeptide comprising a specific domain or truncation of the polypeptides of SEQ ID NO: 445-888. Domains of interest may depend on the nature of the encoded polypeptide; e.g., domains in receptor-like polypeptides include ligand-binding,

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extracellular, transmembrane, or cytoplasmic domains, or combinations thereof, domains in immunoglobulin-like proteins include the variable immunoglobulin-like domains; domains in enzyme-like polypeptides include catalytic and substrate binding domains; and domains in ligand polypeptides include receptor-binding domains.

The polynucleotides of the invention include naturally occurring or wholly or partially synthetic DNA, e.g., cDNA and genomic DNA, and RNA, e.g., mRNA. The polynucleotides may include all of the coding region of the cDNA or may represent a portion of the coding region of the cDNA.

The present invention also provides genes corresponding to the cDNA sequences disclosed herein. The corresponding genes can be isolated in accordance with known methods using the sequence information disclosed herein. Such methods include the preparation of probes or primers from the disclosed sequence information for identification and/or amplification of genes in appropriate genomic libraries or other sources of genomic materials. Further 5' and 3' sequence can be obtained using methods known in the art. For example, full length cDNA or genomic DNA that corresponds to any of the polynucleotides of SEQ ID NO: 1-444 can be obtained by screening appropriate cDNA or genomic DNA libraries under suitable hybridization conditions using any of the polynucleotides of SEQ ID NO: 1-444 or a portion thereof as a probe. Alternatively, the polynucleotides of SEQ ID NO: 1-444 may be used as the basis for suitable primer(s) that allow identification and/or amplification of genes in appropriate genomic DNA or cDNA libraries.

The nucleic acid sequences of the invention can be assembled from ESTs and sequences (including cDNA and genomic sequences) obtained from one or more public databases, such as dbEST, gbpri, and UniGene. The EST sequences can provide identifying sequence information, representative fragment or segment information, or novel segment information for the full-length gene.

The polynucleotides of the invention also provide polynucleotides including nucleotide sequences that are substantially equivalent to the polynucleotides recited above. Polynucleotides according to the invention can have, e.g., at least about 65%, at least about 70%, at least about 75%, at least about 80%, 81%, 82%, 83%, 84%, more typically at least about 85%, 86%, 87%, 88%, 89%, more typically at least about 90%, 91%, 92%, 93%, 94%, and even more typically at least about 95%, 96%, 97%, 98%, 99%, sequence identity to a polynucleotide recited above.

Included within the scope of the nucleic acid sequences of the invention are nucleic acid sequence fragments that hybridize under stringent conditions to any of the nucleotide sequences of SEQ ID NO: 1-444, or complements thereof, which fragment is greater than about 5 nucleotides, preferably 7 nucleotides, more preferably greater than 9 nucleotides and most preferably greater than 17 nucleotides. Fragments of, e.g. 15, 17, or 20 nucleotides or more that are selective for (i.e. specifically hybridize to) any one of the polynucleotides of the invention are contemplated. Probes capable of specifically hybridizing to a polynucleotide can differentiate polynucleotide sequences of the invention from other polynucleotide sequences in the same family of genes or can differentiate human genes from genes of other species, and are preferably based on unique nucleotide sequences.

The sequences falling within the scope of the present invention are not limited to these specific sequences, but also include allelic and species variations thereof. Allelic and species variations can be routinely determined by comparing the sequence provided in SEQ ID NO: 1-444, a representative fragment thereof, or a nucleotide sequence at least 90% identical, preferably 95% identical, to SEQ ID NO: 1-444 with a sequence from another isolate of the same species. Furthermore, to accommodate codon variability, the invention includes nucleic acid molecules coding for the same amino acid sequences as do the specific ORFs disclosed herein. In other words, in the coding region of an ORF, substitution of one codon for another codon that encodes the same amino acid is expressly contemplated.

The nearest neighbor or homology result for the nucleic acids of the present invention, including SEQ ID NO: 1-444, can be obtained by searching a database using an algorithm or a program. Preferably, a BLAST which stands for Basic Local Alignment Search Tool is used to search for local sequence alignments (Altshul, S.F. J Mol. Evol. 36 290-300 (1993) and Altschul S.F. et al. J. Mol. Biol. 21:403-410 (1990)). Alternatively a FASTA version 3 search against Genpept, using Fastxy algorithm.

Species homologs (or orthologs) of the disclosed polynucleotides and proteins are also provided by the present invention. Species homologs may be isolated and identified by making suitable probes or primers from the sequences provided herein and screening a suitable nucleic acid source from the desired species.

The invention also encompasses allelic variants of the disclosed polynucleotides or proteins; that is, naturally-occurring alternative forms of the isolated polynucleotide which also encode proteins which are identical, homologous or related to that encoded by the polynucleotides.

The nucleic acid sequences of the invention are further directed to sequences which encode variants of the described nucleic acids. These amino acid sequence variants may be prepared by methods known in the art by introducing appropriate nucleotide changes into a native or variant polynucleotide. There are two variables in the construction of amino acid sequence variants: the location of the mutation and the nature of the mutation. Nucleic acids encoding the amino acid sequence variants are preferably constructed by mutating the polynucleotide to encode an amino acid sequence that does not occur in nature. These nucleic acid alterations can be made at sites that differ in the nucleic acids from different species (variable positions) or in highly conserved regions (constant regions). Sites at such locations will typically be modified in series, e.g., by substituting first with conservative choices (e.g., hydrophobic amino acid to a different hydrophobic amino acid) and then with more distant choices (e.g., hydrophobic amino acid to a charged amino acid), and then deletions or insertions may be made at the target site. Amino acid sequence deletions generally range from about 1 to 30 residues, preferably about 1 to 10 residues, and are typically contiguous. Amino acid insertions include amino- and/or carboxyl-terminal fusions ranging in length from one to one hundred or more residues, as well as intrasequence insertions of single or multiple amino acid residues. Intrasequence insertions may range generally from about 1 to 10 amino residues, preferably from 1 to 5 residues. Examples of terminal insertions include the heterologous signal sequences necessary for secretion or for intracellular targeting in different host cells and sequences such as FLAG or poly-histidine sequences useful for purifying the expressed protein.

In a preferred method, polynucleotides encoding the novel amino acid sequences are changed via site-directed mutagenesis. This method uses oligonucleotide sequences to alter a polynucleotide to encode the desired amino acid variant, as well as sufficient adjacent nucleotides on both sides of the changed amino acid to form a stable duplex on either side of the site of being changed. In general, the techniques of site-directed mutagenesis are well known to those of skill in the art and this technique is exemplified by publications such as, Edelman et al., DNA 2:183 (1983). A versatile and efficient method for producing site-specific changes in a polynucleotide sequence was published by Zoller and Smith, Nucleic Acids Res. 10:6487-6500 (1982). PCR may also be used to create amino acid sequence variants of the novel nucleic acids. When small amounts of template DNA are used as starting material, primer(s) that differs slightly in sequence from the corresponding region in the template DNA can generate the desired amino acid variant. PCR amplification results in a population of product DNA fragments that differ from the polynucleotide template encoding the polypeptide at the position specified by the primer. The product DNA fragments replace the corresponding region in the plasmid and this gives a polynucleotide encoding the desired amino acid variant.

A further technique for generating amino acid variants is the cassette mutagenesis technique described in Wells et al., *Gene* 34:315 (1985); and other mutagenesis techniques well known in the art, such as, for example, the techniques in Sambrook et al., supra, and *Current Protocols in Molecular Biology*, Ausubel et al. Due to the inherent degeneracy of the genetic code, other DNA sequences which encode substantially the same or a functionally equivalent amino acid sequence may be used in the practice of the invention for the cloning and expression

of these novel nucleic acids. Such DNA sequences include those which are capable of hybridizing to the appropriate novel nucleic acid sequence under stringent conditions.

Polynucleotides encoding preferred polypeptide truncations of the invention can be used to generate polynucleotides encoding chimeric or fusion proteins comprising one or more domains of the invention and heterologous protein sequences.

The polynucleotides of the invention additionally include the complement of any of the polynucleotides recited above. The polynucleotide can be DNA (genomic, cDNA, amplified, or synthetic) or RNA. Methods and algorithms for obtaining such polynucleotides are well known to those of skill in the art and can include, for example, methods for determining hybridization conditions that can routinely isolate polynucleotides of the desired sequence identities.

In accordance with the invention, polynucleotide sequences comprising the mature protein coding sequences corresponding to any one of SEQ ID NO: 1-444, or functional equivalents thereof, may be used to generate recombinant DNA molecules that direct the expression of that nucleic acid, or a functional equivalent thereof, in appropriate host cells. Also included are the cDNA inserts of any of the clones identified herein.

A polynucleotide according to the invention can be joined to any of a variety of other nucleotide sequences by well-established recombinant DNA techniques (see Sambrook J et al. (1989) Molecular Cloning: A Laboratory Manual, Cold Spring Harbor Laboratory, NY). Useful nucleotide sequences for joining to polynucleotides include an assortment of vectors, e.g., plasmids, cosmids, lambda phage derivatives, phagemids, and the like, that are well known in the art. Accordingly, the invention also provides a vector including a polynucleotide of the invention and a host cell containing the polynucleotide. In general, the vector contains an origin of replication functional in at least one organism, convenient restriction endonuclease sites, and a selectable marker for the host cell. Vectors according to the invention include expression vectors, replication vectors, probe generation vectors, and sequencing vectors. A host cell according to the invention can be a prokaryotic or eukaryotic cell and can be a unicellular organism or part of a multicellular organism.

The present invention further provides recombinant constructs comprising a nucleic acid having any of the nucleotide sequences of SEQ ID NO: 1-444 or a fragment thereof or any other polynucleotides of the invention. In one embodiment, the recombinant constructs of the present invention comprise a vector, such as a plasmid or viral vector, into which a nucleic acid having any of the nucleotide sequences of SEQ ID NO: 1-444 or a fragment thereof is inserted, in a forward or reverse orientation. In the case of a vector comprising one of the ORFs of the present invention, the vector may further comprise regulatory sequences, including for example, a promoter, operably linked to the ORF. Large numbers of suitable vectors and promoters are

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known to those of skill in the art and are commercially available for generating the recombinant constructs of the present invention. The following vectors are provided by way of example. Bacterial: pBs, phagescript, PsiX174, pBluescript SK, pBs KS, pNH8a, pNH16a, pNH18a, pNH46a (Stratagene); pTrc99A, pKK223-3, pKK233-3, pDR540, pRIT5 (Pharmacia). Eukaryotic: pWLneo, pSV2cat, pOG44, PXTI, pSG (Stratagene) pSVK3, pBPV, pMSG, pSVL (Pharmacia).

The isolated polynucleotide of the invention may be operably linked to an expression control sequence such as the pMT2 or pED expression vectors disclosed in Kaufman et al., Nucleic Acids Res. 19, 4485-4490 (1991), in order to produce the protein recombinantly. Many suitable expression control sequences are known in the art. General methods of expressing recombinant proteins are also known and are exemplified in R. Kaufman, Methods in Enzymology 185, 537-566 (1990). As defined herein "operably linked" means that the isolated polynucleotide of the invention and an expression control sequence are situated within a vector or cell in such a way that the protein is expressed by a host cell which has been transformed (transfected) with the ligated polynucleotide/expression control sequence.

Promoter regions can be selected from any desired gene using CAT (chloramphenicol transferase) vectors or other vectors with selectable markers. Two appropriate vectors are pKK232-8 and pCM7. Particular named bacterial promoters include lacI, lacZ, T3, T7, gpt, lambda PR, and trc. Eukaryotic promoters include CMV immediate early, HSV thymidine kinase, early and late SV40, LTRs from retrovirus, and mouse metallothionein-I. Selection of the appropriate vector and promoter is well within the level of ordinary skill in the art. Generally, recombinant expression vectors will include origins of replication and selectable markers permitting transformation of the host cell, e.g., the ampicillin resistance gene of E. coli and S. cerevisiae TRP1 gene, and a promoter derived from a highly-expressed gene to direct transcription of a downstream structural sequence. Such promoters can be derived from operons encoding glycolytic enzymes such as 3-phosphoglycerate kinase (PGK), a-factor, acid phosphatase, or heat shock proteins, among others. The heterologous structural sequence is assembled in appropriate phase with translation initiation and termination sequences, and preferably, a leader sequence capable of directing secretion of translated protein into the periplasmic space or extracellular medium. Optionally, the heterologous sequence can encode a fusion protein including an amino terminal identification peptide imparting desired characteristics, e.g., stabilization or simplified purification of expressed recombinant product. Useful expression vectors for bacterial use are constructed by inserting a structural DNA sequence encoding a desired protein together with suitable translation initiation and termination signals in operable reading phase with a functional promoter. The vector will comprise one or

more phenotypic selectable markers and an origin of replication to ensure maintenance of the vector and to, if desirable, provide amplification within the host. Suitable prokaryotic hosts for transformation include *E. coli*, *Bacillus subtilis*, *Salmonella typhimurium* and various species within the genera *Pseudomonas*, *Streptomyces*, and *Staphylococcus*, although others may also be employed as a matter of choice.

As a representative but non-limiting example, useful expression vectors for bacterial use can comprise a selectable marker and bacterial origin of replication derived from commercially available plasmids comprising genetic elements of the well known cloning vector pBR322 (ATCC 37017). Such commercial vectors include, for example, pKK223-3 (Pharmacia Fine Chemicals, Uppsala, Sweden) and GEM 1 (Promega Biotech, Madison, WI, USA). These pBR322 "backbone" sections are combined with an appropriate promoter and the structural sequence to be expressed. Following transformation of a suitable host strain and growth of the host strain to an appropriate cell density, the selected promoter is induced or derepressed by appropriate means (e.g., temperature shift or chemical induction) and cells are cultured for an additional period. Cells are typically harvested by centrifugation, disrupted by physical or chemical means, and the resulting crude extract retained for further purification.

Polynucleotides of the invention can also be used to induce immune responses. For example, as described in Fan et al., *Nat. Biotech.* 17:870-872 (1999), incorporated herein by reference, nucleic acid sequences encoding a polypeptide may be used to generate antibodies against the encoded polypeptide following topical administration of naked plasmid DNA or following injection, and preferably intramuscular injection of the DNA. The nucleic acid sequences are preferably inserted in a recombinant expression vector and may be in the form of naked DNA.

4.3 ANTISENSE NUCLEIC ACIDS

Another aspect of the invention pertains to isolated antisense nucleic acid molecules that are hybridizable to or complementary to the nucleic acid molecule comprising the nucleotide sequence of SEQ ID NO: 1-444, or fragments, analogs or derivatives thereof. An "antisense" nucleic acid comprises a nucleotide sequence that is complementary to a "sense" nucleic acid encoding a protein, e.g., complementary to the coding strand of a double-stranded cDNA molecule or complementary to an mRNA sequence. In specific aspects, antisense nucleic acid molecules are provided that comprise a sequence complementary to at least about 10, 25, 50, 100, 250 or 500 nucleotides or an entire coding strand, or to only a portion thereof. Nucleic acid molecules encoding fragments, homologs, derivatives and analogs of a protein of any of SEQ ID

NO: 445-888 or antisense nucleic acids complementary to a nucleic acid sequence of SEQ ID NO: 1-444 are additionally provided.

In one embodiment, an antisense nucleic acid molecule is antisense to a "coding region" of the coding strand of a nucleotide sequence of the invention. The term "coding region" refers to the region of the nucleotide sequence comprising codons which are translated into amino acid residues. In another embodiment, the antisense nucleic acid molecule is antisense to a "noncoding region" of the coding strand of a nucleotide sequence of the invention. The term "noncoding region" refers to 5' and 3' sequences which flank the coding region that are not translated into amino acids (i.e., also referred to as 5' and 3' untranslated regions).

Given the coding strand sequences encoding a nucleic acid disclosed herein (e.g., SEQ ID NO: 1-444), antisense nucleic acids of the invention can be designed according to the rules of Watson and Crick or Hoogsteen base pairing. The antisense nucleic acid molecule can be complementary to the entire coding region of an mRNA, but more preferably is an oligonucleotide that is antisense to only a portion of the coding or noncoding region of a mRNA. For example, the antisense oligonucleotide can be complementary to the region surrounding the translation start site of a mRNA. An antisense oligonucleotide can be, for example, about 5, 10, 15, 20, 25, 30, 35, 40, 45 or 50 nucleotides in length. An antisense nucleic acid of the invention can be constructed using chemical synthesis or enzymatic ligation reactions using procedures known in the art. For example, an antisense nucleic acid (e.g., an antisense oligonucleotide) can be chemically synthesized using naturally occurring nucleotides or variously modified nucleotides designed to increase the biological stability of the molecules or to increase the physical stability of the duplex formed between the antisense and sense nucleic acids, e.g., phosphorothioate derivatives and acridine substituted nucleotides can be used.

Examples of modified nucleotides that can be used to generate the antisense nucleic acid include: 5-fluorouracil, 5-bromouracil, 5-chlorouracil, 5-iodouracil, hypoxanthine, xanthine, 4-acetylcytosine, 5-(carboxyhydroxylmethyl) uracil, 5-carboxymethylaminomethyl-2-thiouridine, 5-carboxymethylaminomethyluracil, dihydrouracil, beta-D-galactosylqueosine, inosine, N6-isopentenyladenine, 1-methylguanine, 1-methylinosine, 2,2-dimethylguanine, 2-methyladenine, 2-methylguanine, 3-methylcytosine, 5-methylcytosine, N6-adenine, 7-methylguanine, 5-methylaminomethyluracil, 5-methoxyaminomethyl-2-thiouracil, beta-D-mannosylqueosine, 5'-methoxycarboxymethyluracil, 5-methoxyuracil, 2-methylthio-N6-isopentenyladenine, uracil-5-oxyacetic acid (v), wybutoxosine, pseudouracil, queosine, 2-thiocytosine, 5-methyl-2-thiouracil, 2-thiouracil, 4-thiouracil, 5-methyluracil, uracil-5-oxyacetic acid methylester, uracil-5-oxyacetic acid (v), 5-methyl-2-thiouracil, 3-(3-amino-3-N-2-carboxypropyl) uracil, (acp3)w, and 2,6-diaminopurine. Alternatively, the

antisense nucleic acid can be produced biologically using an expression vector into which a nucleic acid has been subcloned in an antisense orientation (i.e., RNA transcribed from the inserted nucleic acid will be of an antisense orientation to a target nucleic acid of interest, described further in the following subsection).

The antisense nucleic acid molecules of the invention are typically administered to a subject or generated in situ such that they hybridize with or bind to cellular mRNA and/or genomic DNA encoding a protein according to the invention to thereby inhibit expression of the protein, e.g., by inhibiting transcription and/or translation. The hybridization can be by conventional nucleotide complementarity to form a stable duplex, or, for example, in the case of an antisense nucleic acid molecule that binds to DNA duplexes, through specific interactions in the major groove of the double helix. An example of a route of administration of antisense nucleic acid molecules of the invention includes direct injection at a tissue site. Alternatively, antisense nucleic acid molecules can be modified to target selected cells and then administered systemically. For example, for systemic administration, antisense molecules can be modified such that they specifically bind to receptors or antigens expressed on a selected cell surface, e.g., by linking the antisense nucleic acid molecules to peptides or antibodies that bind to cell surface receptors or antigens. The antisense nucleic acid molecules can also be delivered to cells using the vectors described herein. To achieve sufficient intracellular concentrations of antisense molecules, vector constructs in which the antisense nucleic acid molecule is placed under the control of a strong pol II or pol III promoter are preferred.

In yet another embodiment, the antisense nucleic acid molecule of the invention is an α-anomeric nucleic acid molecule. An α-anomeric nucleic acid molecule forms specific double-stranded hybrids with complementary RNA in which, contrary to the usual β-units, the strands run parallel to each other (Gaultier et al. (1987) Nucleic Acids Res 15: 6625-6641). The antisense nucleic acid molecule can also comprise a 2'-o-methylribonucleotide (Inoue et al. (1987) Nucleic Acids Res 15: 6131-6148) or a chimeric RNA -DNA analogue (Inoue et al. (1987) FEBS Lett 215: 327-330).

4.4 RIBOZYMES AND PNA MOIETIES

In still another embodiment, an antisense nucleic acid of the invention is a ribozyme. Ribozymes are catalytic RNA molecules with ribonuclease activity that are capable of cleaving a single-stranded nucleic acid, such as an mRNA, to which they have a complementary region. Thus, ribozymes (e.g., hammerhead ribozymes (described in Haselhoff and Gerlach (1988) Nature 334:585-591)) can be used to catalytically cleave a mRNA transcripts to thereby inhibit translation of a mRNA. A ribozyme having specificity for a nucleic acid of the invention can be

designed based upon the nucleotide sequence of a DNA disclosed herein (i.e., SEQ ID NO: 1244). For example, a derivative of a Tetrahymena L-19 IVS RNA can be constructed in which the nucleotide sequence of the active site is complementary to the nucleotide sequence to be cleaved in an mRNA of SEQ ID NO: 1-444 (see, e.g., Cech et al. U.S. Pat. No. 4,987,071; and Cech et al. U.S. Pat. No. 5,116,742). Alternatively, polynucleotides of the invention can be used to select a catalytic RNA having a specific ribonuclease activity from a pool of RNA molecules. See, e.g., Bartel et al., (1993) Science 261:1411-1418.

Alternatively, gene expression can be inhibited by targeting nucleotide sequences complementary to the regulatory region (e.g., promoter and/or enhancers) to form triple helical structures that prevent transcription of the gene in target cells. See generally, Helene. (1991)

Anticancer Drug Des. 6: 569-84; Helene. et al. (1992) Ann. N.Y. Acad. Sci. 660:27-36; and Maher (1992) Bioassays 14: 807-15.

In various embodiments, the nucleic acids of the invention can be modified at the base moiety, sugar moiety or phosphate backbone to improve, e.g., the stability, hybridization, or solubility of the molecule. For example, the deoxyribose phosphate backbone of the nucleic acids can be modified to generate peptide nucleic acids (see Hyrup et al. (1996) Bioorg Med Chem 4: 5-23). As used herein, the terms "peptide nucleic acids" or "PNAs" refer to nucleic acid mimics, e.g., DNA mimics, in which the deoxyribose phosphate backbone is replaced by a pseudopeptide backbone and only the four natural nucleobases are retained. The neutral backbone of PNAs has been shown to allow for specific hybridization to DNA and RNA under conditions of low ionic strength. The synthesis of PNA oligomers can be performed using standard solid phase peptide synthesis protocols as described in Hyrup et al. (1996) above; Perry-O'Keefe et al. (1996) PNAS 93: 14670-675.

PNAs of the invention can be used in therapeutic and diagnostic applications. For example, PNAs can be used as antisense or antigene agents for sequence-specific modulation of gene expression by, e.g., inducing transcription or translation arrest or inhibiting replication. PNAs of the invention can also be used, e.g., in the analysis of single base pair mutations in a gene by, e.g., PNA directed PCR clamping; as artificial restriction enzymes when used in combination with other enzymes, e.g., S1 nucleases (Hyrup B. (1996) above); or as probes or primers for DNA sequence and hybridization (Hyrup et al. (1996), above; Perry-O'Keefe (1996), above).

In another embodiment, PNAs of the invention can be modified, e.g., to enhance their stability or cellular uptake, by attaching lipophilic or other helper groups to PNA, by the formation of PNA-DNA chimeras, or by the use of liposomes or other techniques of drug delivery known in the art. For example, PNA-DNA chimeras can be generated that may

combine the advantageous properties of PNA and DNA. Such chimeras allow DNA recognition enzymes, e.g., RNase H and DNA polymerases, to interact with the DNA portion while the PNA portion would provide high binding affinity and specificity. PNA-DNA chimeras can be linked using linkers of appropriate lengths selected in terms of base stacking, number of bonds between the nucleobases, and orientation (Hyrup (1996) above). The synthesis of PNA-DNA chimeras can be performed as described in Hyrup (1996) above and Finn et al. (1996) Nucl Acids Res 24: 3357-63. For example, a DNA chain can be synthesized on a solid support using standard phosphoramidite coupling chemistry, and modified nucleoside analogs, e.g., 5'-(4-methoxytrityl)amino-5'-deoxy-thymidine phosphoramidite, can be used between the PNA and the 5' end of DNA (Mag et al. (1989) Nucl Acid Res 17: 5973-88). PNA monomers are then coupled in a stepwise manner to produce a chimeric molecule with a 5' PNA segment and a 3' DNA segment (Finn et al. (1996) above). Alternatively, chimeric molecules can be synthesized with a 5' DNA segment and a 3' PNA segment. See, Petersen et al. (1975) Bioorg Med Chem Lett 5: 1119-11124.

In other embodiments, the oligonucleotide may include other appended groups such as peptides (e.g., for targeting host cell receptors in vivo), or agents facilitating transport across the cell membrane (see, e.g., Letsinger et al., 1989, Proc. Natl. Acad. Sci. U.S.A. 86:6553-6556; Lemaitre et al., 1987, Proc. Natl. Acad. Sci. 84:648-652; PCT Publication No. W088/09810) or the blood-brain barrier (see, e.g., PCT Publication No. W089/10134). In addition, oligonucleotides can be modified with hybridization triggered cleavage agents (See, e.g., Krol et al., 1988, BioTechniques 6:958-976) or intercalating agents. (See, e.g., Zon, 1988, Pharm. Res. 5: 539-549). To this end, the oligonucleotide may be conjugated to another molecule, e.g., a peptide, a hybridization triggered cross-linking agent, a transport agent, a hybridization-triggered cleavage agent, etc.

4.5 HOSTS

The present invention further provides host cells genetically engineered to contain the polynucleotides of the invention. For example, such host cells may contain nucleic acids of the invention introduced into the host cell using known transformation, transfection or infection methods. The present invention still further provides host cells genetically engineered to express the polynucleotides of the invention, wherein such polynucleotides are in operative association with a regulatory sequence heterologous to the host cell which drives expression of the polynucleotides in the cell.

Knowledge of nucleic acid sequences allows for modification of cells to permit, or increase, expression of endogenous polypeptide. Cells can be modified (e.g., by homologous

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recombination) to provide increased polypeptide expression by replacing, in whole or in part, the naturally occurring promoter with all or part of a heterologous promoter so that the cells express the polypeptide at higher levels. The heterologous promoter is inserted in such a manner that it is operatively linked to the encoding sequences. See, for example, PCT International Publication No. WO94/12650, PCT International Publication No. WO92/20808, and PCT International Publication No. WO91/09955. It is also contemplated that, in addition to heterologous promoter DNA, amplifiable marker DNA (e.g., ada, dhfr, and the multifunctional CAD gene which encodes carbamyl phosphate synthase, aspartate transcarbamylase, and dihydroorotase) and/or intron DNA may be inserted along with the heterologous promoter DNA. If linked to the coding sequence, amplification of the marker DNA by standard selection methods results in coamplification of the desired protein coding sequences in the cells.

The host cell can be a higher eukaryotic host cell, such as a mammalian cell, a lower eukaryotic host cell, such as a yeast cell, or the host cell can be a prokaryotic cell, such as a bacterial cell. Introduction of the recombinant construct into the host cell can be effected by calcium phosphate transfection, DEAE-dextran mediated transfection, or electroporation (Davis, L. et al., Basic Methods in Molecular Biology (1986)). The host cells containing one of the polynucleotides of the invention, can be used in conventional manners to produce the gene product encoded by the isolated fragment (in the case of an ORF) or can be used to produce a heterologous protein under the control of the EMF.

Any host/vector system can be used to express one or more of the ORFs of the present invention. These include, but are not limited to, eukaryotic hosts such as HeLa cells, Cv-1 cell, COS cells, 293 cells, and Sf9 cells, as well as prokaryotic host such as E. coli and B. suhtilis. The most preferred cells are those which do not normally express the particular polypeptide or protein or which expresses the polypeptide or protein at low natural level. Mature proteins can be expressed in mammalian cells, yeast, bacteria, or other cells under the control of appropriate promoters. Cell-free translation systems can also be employed to produce such proteins using RNAs derived from the DNA constructs of the present invention. Appropriate cloning and expression vectors for use with prokaryotic and eukaryotic hosts are described by Sambrook, et al., in Molecular Cloning: A Laboratory Manual, Second Edition, Cold Spring Harbor, New York (1989), the disclosure of which is hereby incorporated by reference.

Various mammalian cell culture systems can also be employed to express recombinant protein. Examples of mammalian expression systems include the COS-7 lines of monkey kidney fibroblasts, described by Gluzman, Cell 23:175 (1981). Other cell lines capable of expressing a compatible vector are, for example, the C127, monkey COS cells, Chinese Hamster Ovary (CHO) cells, human kidney 293 cells, human epidermal A431 cells, human Colo205 cells, 3T3

cells, CV-1 cells, other transformed primate cell lines, normal diploid cells, cell strains derived from *in vitro* culture of primary tissue, primary explants, HeLa cells, mouse L cells, BHK, HL-60, U937, HaK or Jurkat cells. Mammalian expression vectors will comprise an origin of replication, a suitable promoter and also any necessary ribosome binding sites, polyadenylation site, splice donor and acceptor sites, transcriptional termination sequences, and 5' flanking nontranscribed sequences. DNA sequences derived from the SV40 viral genome, for example, SV40 origin, early promoter, enhancer, splice, and polyadenylation sites may be used to provide the required nontranscribed genetic elements. Recombinant polypeptides and proteins produced in bacterial culture are usually isolated by initial extraction from cell pellets, followed by one or more salting-out, aqueous ion exchange or size exclusion chromatography steps. Protein refolding steps can be used, as necessary, in completing configuration of the mature protein. Finally, high performance liquid chromatography (HPLC) can be employed for final purification steps. Microbial cells employed in expression of proteins can be disrupted by any convenient method, including freeze-thaw cycling, sonication, mechanical disruption, or use of cell lysing agents.

Alternatively, it may be possible to produce the protein in lower eukaryotes such as yeast or insects or in prokaryotes such as bacteria. Potentially suitable yeast strains include Saccharomyces cerevisiae, Schizosaccharomyces pombe, Kluyveromyces strains, Candida, or any yeast strain capable of expressing heterologous proteins. Potentially suitable bacterial strains include Escherichia coli, Bacillus subtilis, Salmonella typhimurium, or any bacterial strain capable of expressing heterologous proteins. If the protein is made in yeast or bacteria, it may be necessary to modify the protein produced therein, for example by phosphorylation or glycosylation of the appropriate sites, in order to obtain the functional protein. Such covalent attachments may be accomplished using known chemical or enzymatic methods.

In another embodiment of the present invention, cells and tissues may be engineered to express an endogenous gene comprising the polynucleotides of the invention under the control of inducible regulatory elements, in which case the regulatory sequences of the endogenous gene may be replaced by homologous recombination. As described herein, gene targeting can be used to replace a gene's existing regulatory region with a regulatory sequence isolated from a different gene or a novel regulatory sequence synthesized by genetic engineering methods. Such regulatory sequences may be comprised of promoters, enhancers, scaffold-attachment regions, negative regulatory elements, transcriptional initiation sites, regulatory protein binding sites or combinations of said sequences. Alternatively, sequences which affect the structure or stability of the RNA or protein produced may be replaced, removed, added, or otherwise modified by targeting. These sequence include polyadenylation signals, mRNA stability elements, splice

sites, leader sequences for enhancing or modifying transport or secretion properties of the protein, or other sequences which alter or improve the function or stability of protein or RNA molecules.

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The targeting event may be a simple insertion of the regulatory sequence, placing the gene under the control of the new regulatory sequence, e.g., inserting a new promoter or enhancer or both upstream of a gene. Alternatively, the targeting event may be a simple deletion of a regulatory element, such as the deletion of a tissue-specific negative regulatory element. Alternatively, the targeting event may replace an existing element; for example, a tissue-specific enhancer can be replaced by an enhancer that has broader or different cell-type specificity than the naturally occurring elements. Here, the naturally occurring sequences are deleted and new sequences are added. In all cases, the identification of the targeting event may be facilitated by the use of one or more selectable marker genes that are contiguous with the targeting DNA, allowing for the selection of cells in which the exogenous DNA has integrated into the host cell genome. The identification of the targeting event may also be facilitated by the use of one or more marker genes exhibiting the property of negative selection, such that the negatively selectable marker is linked to the exogenous DNA, but configured such that the negatively selectable marker flanks the targeting sequence, and such that a correct homologous recombination event with sequences in the host cell genome does not result in the stable integration of the negatively selectable marker. Markers useful for this purpose include the Herpes Simplex Virus thymidine kinase (TK) gene or the bacterial xanthine-guanine phosphoribosyl-transferase (gpt) gene.

The gene targeting or gene activation techniques which can be used in accordance with this aspect of the invention are more particularly described in U.S. Patent No. 5,272,071 to Chappel; U.S. Patent No. 5,578,461 to Sherwin et al.; International Application No. PCT/US92/09627 (WO93/09222) by Selden et al.; and International Application No. PCT/US90/06436 (WO91/06667) by Skoultchi et al., each of which is incorporated by reference herein in its entirety.

4.6 POLYPEPTIDES OF THE INVENTION

The isolated polypeptides of the invention include, but are not limited to, a polypeptide comprising: the amino acid sequences set forth as any one of SEQ ID NO: 445-888 or an amino acid sequence encoded by any one of the nucleotide sequences SEQ ID NO: 1-444 or the corresponding full length or mature protein. Polypeptides of the invention also include polypeptides preferably with biological or immunological activity that are encoded by: (a) a polynucleotide having any one of the nucleotide sequences set forth in SEQ ID NO: 1-444 or (b)

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polynucleotides encoding any one of the amino acid sequences set forth as SEQ ID NO: 445-888 or (c) polynucleotides that hybridize to the complement of the polynucleotides of either (a) or (b) under stringent hybridization conditions. The invention also provides biologically active or immunologically active variants of any of the amino acid sequences set forth as SEQ ID NO: 445-888 or the corresponding full length or mature protein; and "substantial equivalents" thereof (e.g., with at least about 65%, at least about 70%, at least about 75%, at least about 80%, at least about 85%, 86%, 87%, 88%, 89%, at least about 90%, 91%, 92%, 93%, 94%, typically at least about 95%, 96%, 97%, more typically at least about 98%, or most typically at least about 99% amino acid identity) that retain biological activity. Polypeptides encoded by allelic variants may have a similar, increased, or decreased activity compared to polypeptides comprising SEQ ID NO: 445-888.

Fragments of the proteins of the present invention which are capable of exhibiting biological activity are also encompassed by the present invention. Fragments of the protein may be in linear form or they may be cyclized using known methods, for example, as described in H. U. Saragovi, et al., Bio/Technology 10, 773-778 (1992) and in R. S. McDowell, et al., J. Amer. Chem. Soc. 114, 9245-9253 (1992), both of which are incorporated herein by reference. Such fragments may be fused to carrier molecules such as immunoglobulins for many purposes, including increasing the valency of protein binding sites.

The present invention also provides both full-length and mature forms (for example, without a signal sequence or precursor sequence) of the disclosed proteins. The protein coding sequence is identified in the sequence listing by translation of the disclosed nucleotide sequences. The mature form of such protein may be obtained by expression of a full-length polynucleotide in a suitable mammalian cell or other host cell. The sequence of the mature form of the protein is also determinable from the amino acid sequence of the full-length form. Where proteins of the present invention are membrane bound, soluble forms of the proteins are also provided. In such forms, part or all of the regions causing the proteins to be membrane bound are deleted so that the proteins are fully secreted from the cell in which they are expressed.

Protein compositions of the present invention may further comprise an acceptable carrier, such as a hydrophilic, e.g., pharmaceutically acceptable, carrier.

The present invention further provides isolated polypeptides encoded by the nucleic acid fragments of the present invention or by degenerate variants of the nucleic acid fragments of the present invention. By "degenerate variant" is intended nucleotide fragments which differ from a nucleic acid fragment of the present invention (e.g., an ORF) by nucleotide sequence but, due to the degeneracy of the genetic code, encode an identical polypeptide sequence. Preferred nucleic acid fragments of the present invention are the ORFs that encode proteins.

A variety of methodologies known in the art can be utilized to obtain any one of the isolated polypeptides or proteins of the present invention. At the simplest level, the amino acid sequence can be synthesized using commercially available peptide synthesizers. The synthetically-constructed protein sequences, by virtue of sharing primary, secondary or tertiary structural and/or conformational characteristics with proteins may possess biological properties in common therewith, including protein activity. This technique is particularly useful in producing small peptides and fragments of larger polypeptides. Fragments are useful, for example, in generating antibodies against the native polypeptide. Thus, they may be employed as biologically active or immunological substitutes for natural, purified proteins in screening of therapeutic compounds and in immunological processes for the development of antibodies.

The polypeptides and proteins of the present invention can alternatively be purified from cells which have been altered to express the desired polypeptide or protein. As used herein, a cell is said to be altered to express a desired polypeptide or protein when the cell, through genetic manipulation, is made to produce a polypeptide or protein which it normally does not produce or which the cell normally produces at a lower level. One skilled in the art can readily adapt procedures for introducing and expressing either recombinant or synthetic sequences into eukaryotic or prokaryotic cells in order to generate a cell which produces one of the polypeptides or proteins of the present invention.

The invention also relates to methods for producing a polypeptide comprising growing a culture of host cells of the invention in a suitable culture medium, and purifying the protein from the cells or the culture in which the cells are grown. For example, the methods of the invention include a process for producing a polypeptide in which a host cell containing a suitable expression vector that includes a polynucleotide of the invention is cultured under conditions that allow expression of the encoded polypeptide. The polypeptide can be recovered from the culture, conveniently from the culture medium, or from a lysate prepared from the host cells and further purified. Preferred embodiments include those in which the protein produced by such process is a full length or mature form of the protein.

In an alternative method, the polypeptide or protein is purified from bacterial cells which naturally produce the polypeptide or protein. One skilled in the art can readily follow known methods for isolating polypeptides and proteins in order to obtain one of the isolated polypeptides or proteins of the present invention. These include, but are not limited to, immunochromatography, HPLC, size-exclusion chromatography, ion-exchange chromatography, and immuno-affinity chromatography. See, e.g., Scopes, Protein Purification: Principles and Practice, Springer-Verlag (1994); Sambrook, et al., in Molecular Cloning: A Laboratory Manual; Ausubel et al., Current Protocols in Molecular Biology. Polypeptide fragments that

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retain biological/immunological activity include fragments comprising greater than about 100 amino acids, or greater than about 200 amino acids, and fragments that encode specific protein domains.

The purified polypeptides can be used in in vitro binding assays which are well known in the art to identify molecules which bind to the polypeptides. These molecules include but are not limited to, for e.g., small molecules, molecules from combinatorial libraries, antibodies or other proteins. The molecules identified in the binding assay are then tested for antagonist or agonist activity in in vivo tissue culture or animal models that are well known in the art. In brief, the molecules are titrated into a plurality of cell cultures or animals and then tested for either cell/animal death or prolonged survival of the animal/cells.

In addition, the peptides of the invention or molecules capable of binding to the peptides may be complexed with toxins, e.g., ricin or cholera, or with other compounds that are toxic to cells. The toxin-binding molecule complex is then targeted to a tumor or other cell by the specificity of the binding molecule for SEQ ID NO: 445-888.

The protein of the invention may also be expressed as a product of transgenic animals, e.g., as a component of the milk of transgenic cows, goats, pigs, or sheep which are characterized by somatic or germ cells containing a nucleotide sequence encoding the protein.

The proteins provided herein also include proteins characterized by amino acid sequences similar to those of purified proteins but into which modification are naturally provided or deliberately engineered. For example, modifications, in the peptide or DNA sequence, can be made by those skilled in the art using known techniques. Modifications of interest in the protein sequences may include the alteration, substitution, replacement, insertion or deletion of a selected amino acid residue in the coding sequence. For example, one or more of the cysteine residues may be deleted or replaced with another amino acid to alter the conformation of the molecule. Techniques for such alteration, substitution, replacement, insertion or deletion are well known to those skilled in the art (see, e.g., U.S. Pat. No. 4,518,584). Preferably, such alteration, substitution, replacement, insertion or deletion retains the desired activity of the protein. Regions of the protein that are important for the protein function can be determined by various methods known in the art including the alanine-scanning method which involved systematic substitution of single or strings of amino acids with alanine, followed by testing the resulting alanine-containing variant for biological activity. This type of analysis determines the importance of the substituted amino acid(s) in biological activity. Regions of the protein that are important for protein function may be determined by the eMATRIX program.

Other fragments and derivatives of the sequences of proteins which would be expected to retain protein activity in whole or in part and are useful for screening or other immunological

methodologies may also be easily made by those skilled in the art given the disclosures herein..

Such modifications are encompassed by the present invention.

The protein may also be produced by operably linking the isolated polynucleotide of the invention to suitable control sequences in one or more insect expression vectors, and employing an insect expression system. Materials and methods for baculovirus/insect cell expression systems are commercially available in kit form from, e.g., Invitrogen, San Diego, Calif., U.S.A. (the MaxBatTM kit), and such methods are well known in the art, as described in Summers and Smith, Texas Agricultural Experiment Station Bulletin No. 1555 (1987), incorporated herein by reference. As used herein, an insect cell capable of expressing a polynucleotide of the present invention is "transformed."

The protein of the invention may be prepared by culturing transformed host cells under culture conditions suitable to express the recombinant protein. The resulting expressed protein may then be purified from such culture (i.e., from culture medium or cell extracts) using known purification processes, such as gel filtration and ion exchange chromatography. The purification of the protein may also include an affinity column containing agents which will bind to the protein; one or more column steps over such affinity resins as concanavalin A-agarose, heparin-toyopearlTM or Cibacrom blue 3GA SepharoseTM; one or more steps involving hydrophobic interaction chromatography using such resins as phenyl ether, butyl ether, or propyl ether; or immunoaffinity chromatography.

Alternatively, the protein of the invention may also be expressed in a form which will facilitate purification. For example, it may be expressed as a fusion protein, such as those of maltose binding protein (MBP), glutathione-S-transferase (GST) or thioredoxin (TRX), or as a His tag. Kits for expression and purification of such fusion proteins are commercially available from New England BioLab (Beverly, Mass.), Pharmacia (Piscataway, N.J.) and Invitrogen, respectively. The protein can also be tagged with an epitope and subsequently purified by using a specific antibody directed to such epitope. One such epitope ("FLAG®") is commercially available from Kodak (New Haven, Conn.).

Finally, one or more reverse-phase high performance liquid chromatography (RP-HPLC) steps employing hydrophobic RP-HPLC media, e.g., silica gel having pendant methyl or other aliphatic groups, can be employed to further purify the protein. Some or all of the foregoing purification steps, in various combinations, can also be employed to provide a substantially homogeneous isolated recombinant protein. The protein thus purified is substantially free of other mammalian proteins and is defined in accordance with the present invention as an "isolated protein."

The polypeptides of the invention include analogs (variants). This embraces fragments, as well as peptides in which one or more amino acids has been deleted, inserted, or substituted. Also, analogs of the polypeptides of the invention embrace fusions of the polypeptides or modifications of the polypeptides of the invention, wherein the polypeptide or analog is fused to another moiety or moieties, e.g., targeting moiety or another therapeutic agent. Such analogs may exhibit improved properties such as activity and/or stability. Examples of moieties which may be fused to the polypeptide or an analog include, for example, targeting moieties which provide for the delivery of polypeptide to pancreatic cells, e.g., antibodies to pancreatic cells, antibodies to immune cells such as T-cells, monocytes, dendritic cells, granulocytes, etc., as well as receptor and ligands expressed on pancreatic or immune cells. Other moieties which may be fused to the polypeptide include therapeutic agents which are used for treatment, for example, immunosuppressive drugs such as cyclosporin, SK506, azathioprine, CD3 antibodies and steroids. Also, polypeptides may be fused to immune modulators, and other cytokines such as alpha or beta interferon.

4.6.1 DETERMINING POLYPEPTIDE AND POLYNUCLEOTIDE IDENTITY AND SIMILARITY

Preferred identity and/or similarity are designed to give the largest match between the sequences tested. Methods to determine identity and similarity are codified in computer programs including, but are not limited to, the GCG program package, including GAP (Devereux, J., et al., Nucleic Acids Research 12(1):387 (1984); Genetics Computer Group, University of Wisconsin, Madison, WI), BLASTP, BLASTN, BLASTX, FASTA (Altschul, S.F. et al., J. Molec. Biol. 215:403-410 (1990), PSI-BLAST (Altschul S.F. et al., Nucleic Acids Res. vol. 25, pp. 3389-3402, herein incorporated by reference), eMatrix software (Wu et al., J. Comp. Biol., Vol. 6, pp. 219-235 (1999), herein incorporated by reference), eMotif software (Nevill-Manning et al, ISMB-97, Vol. 4, pp. 202-209, herein incorporated by reference), pFam software (Sonnhammer et al., Nucleic Acids Res., Vol. 26(1), pp. 320-322 (1998), herein incorporated by reference), the GeneAtlas software (Molecular Simulations Inc. (MSI), San Diego, CA) (Sanchez and Sali (1998) Proc. Natl. Acad. Sci., 95, 13597-13602; Kitson DH et al, (2000) "Remote homology detection using structural modeling - an evaluation" Submitted; Fischer and Eisenberg (1996) Protein Sci. 5, 947-955), Neural Network SignalP V1.1 program (from Center for Biological Sequence Analysis, The Technical University of Denmark), and the Kyte-Doolittle hydrophobocity prediction algorithm (J. Mol Biol, 157, pp. 105-31 (1982), incorporated herein by reference). The BLAST programs are publicly available from the National Center for Biotechnology Information (NCBI) and other sources (BLAST Manual,

Altschul, S., et al. NCB NLM NIH Bethesda, MD 20894; Altschul, S., et al., J. Mol. Biol. 215:403-410 (1990).

4.7 CHIMERIC AND FUSION PROTEINS

The invention also provides chimeric or fusion proteins. As used herein, a "chimeric protein" or "fusion protein" comprises a polypeptide of the invention operatively linked to another polypeptide. Within a fusion protein the polypeptide according to the invention can correspond to all or a portion of a protein according to the invention. In one embodiment, a fusion protein comprises at least one biologically active portion of a protein according to the invention. In another embodiment, a fusion protein comprises at least two biologically active portions of a protein according to the invention. Within the fusion protein, the term "operatively linked" is intended to indicate that the polypeptide according to the invention and the other polypeptide are fused in-frame to each other. The polypeptide can be fused to the N-terminus or C-terminus.

For example, in one embodiment a fusion protein comprises a polypeptide according to the invention operably linked to the extracellular domain of a second protein. In another embodiment, the fusion protein is a GST-fusion protein in which the polypeptide sequences of the invention are fused to the C-terminus of the GST (i.e., glutathione S-transferase) sequences.

In another embodiment, the fusion protein is an immunoglobulin fusion protein in which the polypeptide sequences according to the invention comprise one or more domains fused to sequences derived from a member of the immunoglobulin protein family. The immunoglobulin fusion proteins of the invention can be incorporated into pharmaceutical compositions and administered to a subject to inhibit an interaction between a ligand and a protein of the invention on the surface of a cell, to thereby suppress signal transduction in vivo. The immunoglobulin fusion proteins can be used to affect the bioavailability of a cognate ligand. Inhibition of the ligand/protein interaction may be useful therapeutically for both the treatment of proliferative and differentiative disorders, e,g., cancer as well as modulating (e.g., promoting or inhibiting) cell survival. Moreover, the immunoglobulin fusion proteins of the invention can be used as immunogens to produce antibodies in a subject, to purify ligands, and in screening assays to identify molecules that inhibit the interaction of a polypeptide of the invention with a ligand.

A chimeric or fusion protein of the invention can be produced by standard recombinant DNA techniques. For example, DNA fragments coding for the different polypeptide sequences are ligated together in-frame in accordance with conventional techniques, e.g., by employing blunt-ended or stagger-ended termini for ligation, restriction enzyme digestion to provide for appropriate termini, filling-in of cohesive ends as appropriate, alkaline phosphatase treatment to avoid undesirable joining, and enzymatic ligation. In another embodiment, the fusion gene can be synthesized by conventional techniques including automated DNA synthesizers. Alternatively, PCR amplification of gene fragments can be carried out using anchor primers that give rise to complementary overhangs between two consecutive gene fragments that can subsequently be annealed and reamplified to generate a chimeric gene sequence (see, for example, Ausubel et al. (eds.) Current Protocols in Molecular Biology, John Wiley & Sons, 1992). Moreover, many expression vectors are commercially available that already encode a fusion moiety (e.g., a GST polypeptide). A nucleic acid encoding a polypeptide of the invention can be cloned into such an expression vector such that the fusion moiety is linked in-frame to the protein of the invention.

4.8 GENE THERAPY

Mutations in the polynucleotides of the invention may result in loss of normal function of the encoded protein. The invention thus provides gene therapy to restore normal activity of the polypeptides of the invention; or to treat disease states involving polypeptides of the invention. Delivery of a functional gene encoding polypeptides of the invention to appropriate cells is effected ex vivo, in situ, or in vivo by use of vectors, and more particularly viral vectors (e.g., adenovirus, adeno-associated virus, or a retrovirus), or ex vivo by use of physical DNA transfer methods (e.g., liposomes or chemical treatments). See, for example, Anderson, Nature, supplement to vol. 392, no. 6679, pp.25-20 (1998). For additional reviews of gene therapy technology see Friedmann, Science, 244: 1275-1281 (1989); Verma, Scientific American: 68-84 (1990); and Miller, Nature, 357: 455-460 (1992). Introduction of any one of the nucleotides of the present invention or a gene encoding the polypeptides of the present invention can also be accomplished with extrachromosomal substrates (transient expression) or artificial chromosomes (stable expression). Cells may also be cultured ex vivo in the presence of proteins of the present invention in order to proliferate or to produce a desired effect on or activity in such cells. Treated cells can then be introduced in vivo for therapeutic purposes. Alternatively, it is contemplated that in other human disease states, preventing the expression of or inhibiting the activity of polypeptides of the invention will be useful in treating the disease states. It is contemplated that antisense therapy or gene therapy could be applied to negatively regulate the expression of polypeptides of the invention.

Other methods inhibiting expression of a protein include the introduction of antisense molecules to the nucleic acids of the present invention, their complements, or their translated RNA sequences, by methods known in the art. Further, the polypeptides of the present invention can be

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inhibited by using targeted deletion methods, or the insertion of a negative regulatory element such as a silencer, which is tissue specific.

The present invention still further provides cells genetically engineered in vivo to express the polynucleotides of the invention, wherein such polynucleotides are in operative association with a regulatory sequence heterologous to the host cell which drives expression of the polynucleotides in the cell. These methods can be used to increase or decrease the expression of the polynucleotides of the present invention.

Knowledge of DNA sequences provided by the invention allows for modification of cells to permit, increase, or decrease, expression of endogenous polypeptide. Cells can be modified (e.g., by homologous recombination) to provide increased polypeptide expression by replacing, in whole or in part, the naturally occurring promoter with all or part of a heterologous promoter so that the cells express the protein at higher levels. The heterologous promoter is inserted in such a manner that it is operatively linked to the desired protein encoding sequences. See, for example, PCT International Publication No. WO 94/12650, PCT International Publication No. WO 92/20808, and PCT International Publication No. WO 91/09955. It is also contemplated that, in addition to heterologous promoter DNA, amplifiable marker DNA (e.g., ada, dhfr, and the multifunctional CAD gene which encodes carbamyl phosphate synthase, aspartate transcarbamylase, and dihydroorotase) and/or intron DNA may be inserted along with the heterologous promoter DNA. If linked to the desired protein coding sequence, amplification of the marker DNA by standard selection methods results in co-amplification of the desired protein coding sequences in the cells.

In another embodiment of the present invention, cells and tissues may be engineered to express an endogenous gene comprising the polynucleotides of the invention under the control of inducible regulatory elements, in which case the regulatory sequences of the endogenous gene may be replaced by homologous recombination. As described herein, gene targeting can be used to replace a gene's existing regulatory region with a regulatory sequence isolated from a different gene or a novel regulatory sequence synthesized by genetic engineering methods. Such regulatory sequences may be comprised of promoters, enhancers, scaffold-attachment regions, negative regulatory elements, transcriptional initiation sites, regulatory protein binding sites or combinations of said sequences. Alternatively, sequences which affect the structure or stability of the RNA or protein produced may be replaced, removed, added, or otherwise modified by targeting. These sequences include polyadenylation signals, mRNA stability elements, splice sites, leader sequences for enhancing or modifying transport or secretion properties of the protein, or other sequences which alter or improve the function or stability of protein or RNA molecules.

The targeting event may be a simple insertion of the regulatory sequence, placing the gene under the control of the new regulatory sequence, e.g., inserting a new promoter or enhancer or both upstream of a gene. Alternatively, the targeting event may be a simple deletion of a regulatory element, such as the deletion of a tissue-specific negative regulatory element. Alternatively, the targeting event may replace an existing element; for example, a tissue-specific enhancer can be replaced by an enhancer that has broader or different cell-type specificity than the naturally occurring elements. Here, the naturally occurring sequences are deleted and new sequences are added. In all cases, the identification of the targeting event may be facilitated by the use of one or more selectable marker genes that are contiguous with the targeting DNA, allowing for the selection of cells in which the exogenous DNA has integrated into the cell genome. The identification of the targeting event may also be facilitated by the use of one or more marker genes exhibiting the property of negative selection, such that the negatively selectable marker is linked to the exogenous DNA, but configured such that the negatively selectable marker flanks the targeting sequence, and such that a correct homologous recombination event with sequences in the host cell genome does not result in the stable integration of the negatively selectable marker. Markers useful for this purpose include the Herpes Simplex Virus thymidine kinase (TK) gene or the bacterial xanthine-guanine phosphoribosyl-transferase (gpt) gene.

The gene targeting or gene activation techniques which can be used in accordance with this aspect of the invention are more particularly described in U.S. Patent No. 5,272,071 to Chappel; U.S. Patent No. 5,578,461 to Sherwin et al.; International Application No. PCT/US92/09627 (WO93/09222) by Selden et al.; and International Application No. PCT/US90/06436 (WO91/06667) by Skoultchi et al., each of which is incorporated by reference herein in its entirety.

4.9 TRANSGENIC ANIMALS

In preferred methods to determine biological functions of the polypeptides of the invention in vivo, one or more genes provided by the invention are either over expressed or inactivated in the germ line of animals using homologous recombination [Capecchi, Science 244:1288-1292 (1989)]. Animals in which the gene is over expressed, under the regulatory control of exogenous or endogenous promoter elements, are known as transgenic animals. Animals in which an endogenous gene has been inactivated by homologous recombination are referred to as "knockout" animals. Knockout animals, preferably non-human mammals, can be prepared as described in U.S. Patent No. 5,557,032, incorporated herein by reference. Transgenic animals are useful to determine the roles polypeptides of the invention play in biological processes, and preferably in disease states. Transgenic animals are useful as model systems to identify compounds that modulate lipid metabolism. Transgenic animals, preferably non-human mammals, are produced using methods as described in U.S. Patent No 5,489,743 and PCT Publication No. WO94/28122, incorporated herein by reference.

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Transgenic animals can be prepared wherein all or part of a promoter of the polynucleotides of the invention is either activated or inactivated to alter the level of expression of the polypeptides of the invention. Inactivation can be carried out using homologous recombination methods described above. Activation can be achieved by supplementing or even replacing the homologous promoter to provide for increased protein expression. The homologous promoter can be supplemented by insertion of one or more heterologous enhancer elements known to confer promoter activation in a particular tissue.

The polynucleotides of the present invention also make possible the development, through, e.g., homologous recombination or knock out strategies, of animals that fail to express polypeptides of the invention or that express a variant polypeptide. Such animals are useful as models for studying the in vivo activities of polypeptide as well as for studying modulators of the polypeptides of the invention.

In preferred methods to determine biological functions of the polypeptides of the invention in vivo, one or more genes provided by the invention are either over expressed or inactivated in the germ line of animals using homologous recombination [Capecchi, Science 244:1288-1292 (1989)]. Animals in which the gene is over expressed, under the regulatory control of exogenous or endogenous promoter elements, are known as transgenic animals. Animals in which an endogenous gene has been inactivated by homologous recombination are referred to as "knockout" animals. Knockout animals, preferably non-human mammals, can be prepared as described in U.S. Patent No. 5,557,032, incorporated herein by reference. Transgenic animals are useful to determine the roles polypeptides of the invention play in biological processes, and preferably in disease states. Transgenic animals are useful as model systems to identify compounds that modulate lipid metabolism. Transgenic animals, preferably non-human mammals, are produced using methods as described in U.S. Patent No 5,489,743 and PCT Publication No. WO94/28122, incorporated herein by reference.

Transgenic animals can be prepared wherein all or part of the polynucleotides of the invention promoter is either activated or inactivated to alter the level of expression of the polypeptides of the invention. Inactivation can be carried out using homologous recombination methods described above. Activation can be achieved by supplementing or even replacing the homologous promoter to provide for increased protein expression. The homologous promoter can be supplemented by insertion of one or more heterologous enhancer elements known to confer promoter activation in a particular tissue.

4.10 USES AND BIOLOGICAL ACTIVITY

The polynucleotides and proteins of the present invention are expected to exhibit one or more of the uses or biological activities (including those associated with assays cited herein) identified herein. Uses or activities described for proteins of the present invention may be provided by administration or use of such proteins or of polynucleotides encoding such proteins (such as, for example, in gene therapies or vectors suitable for introduction of DNA). The mechanism underlying the particular condition or pathology will dictate whether the polypeptides of the invention, the polynucleotides of the invention or modulators (activators or inhibitors) thereof would be beneficial to the subject in need of treatment. Thus, "therapeutic compositions of the invention" include compositions comprising isolated polynucleotides (including recombinant DNA molecules, cloned genes and degenerate variants thereof) or polypeptides of the invention (including full length protein, mature protein and truncations or domains thereof), or compounds and other substances that modulate the overall activity of the target gene products, either at the level of target gene/protein expression or target protein activity. Such modulators include polypeptides, analogs, (variants), including fragments and fusion proteins, antibodies and other binding proteins; chemical compounds that directly or indirectly activate or inhibit the polypeptides of the invention (identified, e.g., via drug screening assays as described herein); antisense polynucleotides and polynucleotides suitable for triple helix formation; and in particular antibodies or other binding partners that specifically recognize one or more epitopes of the polypeptides of the invention.

The polypeptides of the present invention may likewise be involved in cellular activation or in one of the other physiological pathways described herein.

4.10.1 RESEARCH USES AND UTILITIES

The polynucleotides provided by the present invention can be used by the research community for various purposes. The polynucleotides can be used to express recombinant protein for analysis, characterization or therapeutic use; as markers for tissues in which the corresponding protein is preferentially expressed (either constitutively or at a particular stage of tissue differentiation or development or in disease states); as molecular weight markers on gels; as chromosome markers or tags (when labeled) to identify chromosomes or to map related gene positions; to compare with endogenous DNA sequences in patients to identify potential genetic disorders; as probes to hybridize and thus discover novel, related DNA sequences; as a source of information to derive PCR primers for genetic fingerprinting; as a probe to "subtract-out" known sequences in the process of discovering other novel polynucleotides; for selecting and making oligomers for attachment to a "gene chip" or other support, including for examination of expression patterns; to raise anti-protein antibodies using DNA immunization techniques; and as

an antigen to raise anti-DNA antibodies or elicit another immune response. Where the polynucleotide encodes a protein which binds or potentially binds to another protein (such as, for example, in a receptor-ligand interaction), the polynucleotide can also be used in interaction trap assays (such as, for example, that described in Gyuris et al., Cell 75:791-803 (1993)) to identify polynucleotides encoding the other protein with which binding occurs or to identify inhibitors of the binding interaction.

The polypeptides provided by the present invention can similarly be used in assays to determine biological activity, including in a panel of multiple proteins for high-throughput screening; to raise antibodies or to elicit another immune response; as a reagent (including the labeled reagent) in assays designed to quantitatively determine levels of the protein (or its receptor) in biological fluids; as markers for tissues in which the corresponding polypeptide is preferentially expressed (either constitutively or at a particular stage of tissue differentiation or development or in a disease state); and, of course, to isolate correlative receptors or ligands. Proteins involved in these binding interactions can also be used to screen for peptide or small molecule inhibitors or agonists of the binding interaction.

Any or all of these research utilities are capable of being developed into reagent grade or kit format for commercialization as research products.

Methods for performing the uses listed above are well known to those skilled in the art. References disclosing such methods include without limitation "Molecular Cloning: A Laboratory Manual", 2d ed., Cold Spring Harbor Laboratory Press, Sambrook, J., E. F. Fritsch and T. Maniatis eds., 1989, and "Methods in Enzymology: Guide to Molecular Cloning Techniques", Academic Press, Berger, S. L. and A. R. Kimmel eds., 1987.

4.10.2 NUTRITIONAL USES

Polynucleotides and polypeptides of the present invention can also be used as nutritional sources or supplements. Such uses include without limitation use as a protein or amino acid supplement, use as a carbon source, use as a nitrogen source and use as a source of carbohydrate. In such cases the polypeptide or polynucleotide of the invention can be added to the feed of a particular organism or can be administered as a separate solid or liquid preparation, such as in the form of powder, pills, solutions, suspensions or capsules. In the case of microorganisms, the polypeptide or polynucleotide of the invention can be added to the medium in or on which the microorganism is cultured.

4.10.3 CYTOKINE AND CELL PROLIFERATION/DIFFERENTIATION ACTIVITY

A polypeptide of the present invention may exhibit activity relating to cytokine, cell proliferation (either inducing or inhibiting) or cell differentiation (either inducing or inhibiting) activity or may induce production of other cytokines in certain cell populations. A polynucleotide of the invention can encode a polypeptide exhibiting such attributes. Many protein factors discovered to date, including all known cytokines, have exhibited activity in one or more factor-dependent cell proliferation assays, and hence the assays serve as a convenient confirmation of cytokine activity. The activity of therapeutic compositions of the present invention is evidenced by any one of a number of routine factor dependent cell proliferation assays for cell lines including, without limitation, 32D, DA2, DA1G, T10, B9, B9/11, BaF3, MC9/G, M+(preB M+), 2E8, RB5, DA1, 123, T1165, HT2, CTLL2, TF-1, Mo7e, CMK, HUVEC, and Caco. Therapeutic compositions of the invention can be used in the following:

Assays for T-cell or thymocyte proliferation include without limitation those described in: Current Protocols in Immunology, Ed by J. E. Coligan, A. M. Kruisbeek, D. H. Margulies, E. M. Shevach, W. Strober, Pub. Greene Publishing Associates and Wiley-Interscience (Chapter 3, In Vitro assays for Mouse Lymphocyte Function 3.1-3.19; Chapter 7, Immunologic studies in Humans); Takai et al., J. Immunol. 137:3494-3500, 1986; Bertagnolli et al., J. Immunol. 145:1706-1712, 1990; Bertagnolli et al., Cellular Immunology 133:327-341, 1991; Bertagnolli, et al., I. Immunol. 149:3778-3783, 1992; Bowman et al., I. Immunol. 152:1756-1761, 1994.

Assays for cytokine production and/or proliferation of spleen cells, lymph node cells or thymocytes include, without limitation, those described in: Polyclonal T cell stimulation, Kruisbeek, A. M. and Shevach, E. M. In Current Protocols in Immunology. J. E. e.a. Coligan eds. Vol 1 pp. 3.12.1-3.12.14, John Wiley and Sons, Toronto. 1994; and Measurement of mouse and human interleukin-y, Schreiber, R. D. In Current Protocols in Immunology. J. E. e.a. Coligan eds. Vol 1 pp. 6.8.1-6.8.8, John Wiley and Sons, Toronto. 1994.

Assays for proliferation and differentiation of hematopoietic and lymphopoietic cells include, without limitation, those described in: Measurement of Human and Murine Interleukin 2 and Interleukin 4, Bottomly, K., Davis, L. S. and Lipsky, P. E. In Current Protocols in Immunology, J. E. e.a. Coligan eds. Vol 1 pp. 6.3.1-6.3.12, John Wiley and Sons, Toronto. 1991; deVries et al., J. Exp. Med. 173:1205-1211, 1991; Moreau et al., Nature 336:690-692, 1988; Greenberger et al., Proc. Natl. Acad. Sci. U.S.A. 80:2931-2938, 1983; Measurement of mouse and human interleukin 6--Nordan, R. In Current Protocols in Immunology. J. E. Coligan eds. Vol 1 pp. 6.6.1-6.6.5, John Wiley and Sons, Toronto. 1991; Smith et al., Proc. Natl. Aced. Sci. U.S.A. 83:1857-1861, 1986; Measurement of human Interleukin 11--Bennett, F., Giannotti, J., Clark, S. C. and Turner, K. J. In Current Protocols in Immunology. J. E. Coligan eds. Vol 1 pp. 6.15.1 John Wiley and Sons, Toronto. 1991; Measurement of mouse and human Interleukin

9--Ciarletta, A., Giannotti, J., Clark, S. C. and Turner, K. J. In Current Protocols in Immunology. J. E. Coligan eds. Vol 1 pp. 6.13.1, John Wiley and Sons, Toronto. 1991.

Assays for T-cell clone responses to antigens (which will identify, among others, proteins that affect APC-T cell interactions as well as direct T-cell effects by measuring proliferation and cytokine production) include, without limitation, those described in: Current Protocols in Immunology, Ed by J. E. Coligan, A. M. Kruisbeek, D. H. Margulies, E. M. Shevach, W Strober, Pub. Greene Publishing Associates and Wiley-Interscience (Chapter 3, In Vitro assays for Mouse Lymphocyte Function; Chapter 6, Cytokines and their cellular receptors; Chapter 7, Immunologic studies in Humans); Weinberger et al., Proc. Natl. Acad. Sci. USA 77:6091-6095. 1980; Weinberger et al., Eur. J. Immun. 11:405-411, 1981; Takai et al., J. Immunol. 137:3494-3500, 1986; Takai et al., J. Immunol. 140:508-512, 1988.

4.10.4 STEM CELL GROWTH FACTOR ACTIVITY

A polypeptide of the present invention may exhibit stem cell growth factor activity and be involved in the proliferation, differentiation and survival of pluripotent and totipotent stem cells including primordial germ cells, embryonic stem cells, hematopoietic stem cells and/or germ line stem cells. Administration of the polypeptide of the invention to stem cells in vivo or ex vivo is expected to maintain and expand cell populations in a totipotential or pluripotential state which would be useful for re-engineering damaged or diseased tissues, transplantation, manufacture of bio-pharmaceuticals and the development of bio-sensors. The ability to produce large quantities of human cells has important working applications for the production of human proteins which currently must be obtained from non-human sources or donors, implantation of cells to treat diseases such as Parkinson's, Alzheimer's and other neurodegenerative diseases; tissues for grafting such as bone marrow, skin, cartilage, tendons, bone, muscle (including cardiac muscle), blood vessels, cornea, neural cells, gastrointestinal cells and others; and organs for transplantation such as kidney, liver, pancreas (including islet cells), heart and lung.

It is contemplated that multiple different exogenous growth factors and/or cytokines may be administered in combination with the polypeptide of the invention to achieve the desired effect, including any of the growth factors listed herein, other stem cell maintenance factors, and specifically including stem cell factor (SCF), leukemia inhibitory factor (LIF), Flt-3 ligand (Flt-3L), any of the interleukins, recombinant soluble IL-6 receptor fused to IL-6, macrophage inflammatory protein 1-alpha (MIP-1-alpha), G-CSF, GM-CSF, thrombopoietin (TPO), platelet factor 4 (PF-4), platelet-derived growth factor (PDGF), neural growth factors and basic fibroblast growth factor (bFGF).

Since totipotent stem cells can give rise to virtually any mature cell type, expansion of these cells in culture will facilitate the production of large quantities of mature cells. Techniques for culturing stem cells are known in the art and administration of polypeptides of the invention, optionally with other growth factors and/or cytokines, is expected to enhance the survival and proliferation of the stem cell populations. This can be accomplished by direct administration of the polypeptide of the invention to the culture medium. Alternatively, stroma cells transfected with a polynucleotide that encodes for the polypeptide of the invention can be used as a feeder layer for the stem cell populations in culture or in vivo. Stromal support cells for feeder layers may include embryonic bone marrow fibroblasts, bone marrow stromal cells, fetal liver cells, or cultured embryonic fibroblasts (see U.S. Patent No. 5,690,926).

Stem cells themselves can be transfected with a polynucleotide of the invention to induce autocrine expression of the polypeptide of the invention. This will allow for generation of undifferentiated totipotential/pluripotential stem cell lines that are useful as is or that can then be differentiated into the desired mature cell types. These stable cell lines can also serve as a source of undifferentiated totipotential/pluripotential mRNA to create cDNA libraries and templates for polymerase chain reaction experiments. These studies would allow for the isolation and identification of differentially expressed genes in stem cell populations that regulate stem cell proliferation and/or maintenance.

Expansion and maintenance of totipotent stem cell populations will be useful in the treatment of many pathological conditions. For example, polypeptides of the present invention may be used to manipulate stem cells in culture to give rise to neuroepithelial cells that can be used to augment or replace cells damaged by illness, autoimmune disease, accidental damage or genetic disorders. The polypeptide of the invention may be useful for inducing the proliferation of neural cells and for the regeneration of nerve and brain tissue, i.e. for the treatment of central and peripheral nervous system diseases and neuropathies, as well as mechanical and traumatic disorders which involve degeneration, death or trauma to neural cells or nerve tissue. In addition, the expanded stem cell populations can also be genetically altered for gene therapy purposes and to decrease host rejection of replacement tissues after grafting or implantation.

Expression of the polypeptide of the invention and its effect on stem cells can also be manipulated to achieve controlled differentiation of the stem cells into more differentiated cell types. A broadly applicable method of obtaining pure populations of a specific differentiated cell type from undifferentiated stem cell populations involves the use of a cell-type specific promoter driving a selectable marker. The selectable marker allows only cells of the desired type to survive. For example, stem cells can be induced to differentiate into cardiomyocytes (Wobus et al., Differentiation, 48: 173-182, (1991); Klug et al., J. Clin. Invest., 98(1): 216-224, (1998))

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or skeletal muscle cells (Browder, L. W. In: Principles of Tissue Engineering eds. Lanza et al., Academic Press (1997)). Alternatively, directed differentiation of stem cells can be accomplished by culturing the stem cells in the presence of a differentiation factor such as retinoic acid and an antagonist of the polypeptide of the invention which would inhibit the effects of endogenous stem cell factor activity and allow differentiation to proceed.

In vitro cultures of stem cells can be used to determine if the polypeptide of the invention exhibits stem cell growth factor activity. Stem cells are isolated from any one of various cell sources (including hematopoietic stem cells and embryonic stem cells) and cultured on a feeder layer, as described by Thompson et al. Proc. Natl. Acad. Sci, U.S.A., 92: 7844-7848 (1995), in the presence of the polypeptide of the invention alone or in combination with other growth factors or cytokines. The ability of the polypeptide of the invention to induce stem cells proliferation is determined by colony formation on semi-solid support e.g. as described by Bernstein et al., Blood, 77: 2316-2321 (1991).

4.10.5 HEMATOPOIESIS REGULATING ACTIVITY

A polypeptide of the present invention may be involved in regulation of hematopoiesis and, consequently, in the treatment of myeloid or lymphoid cell disorders. Even marginal biological activity in support of colony forming cells or of factor-dependent cell lines indicates involvement in regulating hematopoiesis, e.g. in supporting the growth and proliferation of erythroid progenitor cells alone or in combination with other cytokines, thereby indicating utility, for example, in treating various anemias or for use in conjunction with irradiation/chemotherapy to stimulate the production of erythroid precursors and/or erythroid cells; in supporting the growth and proliferation of myeloid cells such as granulocytes and monocytes/macrophages (i.e., traditional CSF activity) useful, for example, in conjunction with chemotherapy to prevent or treat consequent myelo-suppression; in supporting the growth and proliferation of megakaryocytes and consequently of platelets thereby allowing prevention or treatment of various platelet disorders such as thrombocytopenia, and generally for use in place of or complimentary to platelet transfusions; and/or in supporting the growth and proliferation of hematopoietic stem cells which are capable of maturing to any and all of the above-mentioned hematopoietic cells and therefore find therapeutic utility in various stem cell disorders (such as those usually treated with transplantation, including, without limitation, aplastic anemia and paroxysmal nocturnal hemoglobinuria), as well as in repopulating the stem cell compartment post irradiation/chemotherapy, either in-vivo or ex-vivo (i.e., in conjunction with bone marrow transplantation or with peripheral progenitor cell transplantation (homologous or heterologous)) as normal cells or genetically manipulated for gene therapy.

Therapeutic compositions of the invention can be used in the following:

Suitable assays for proliferation and differentiation of various hematopoietic lines are cited above.

Assays for embryonic stem cell differentiation (which will identify, among others, proteins that influence embryonic differentiation hematopoiesis) include, without limitation, those described in: Johansson et al. Cellular Biology 15:141-151, 1995; Keller et al., Molecular and Cellular Biology 13:473-486, 1993; McClanahan et al., Blood 81:2903-2915, 1993.

Assays for stem cell survival and differentiation (which will identify, among others, proteins that regulate lympho-hematopoiesis) include, without limitation, those described in: Methylcellulose colony forming assays, Freshney, M. G. In Culture of Hematopoietic Cells. R. I. Freshney, et al. eds. Vol pp. 265-268, Wiley-Liss, Inc., New York, N.Y. 1994; Hirayama et al., Proc. Natl. Acad. Sci. USA 89:5907-5911, 1992; Primitive hematopoietic colony forming cells with high proliferative potential, McNiece, I. K. and Briddell, R. A. In Culture of Hematopoietic Cells. R. I. Freshney, et al. eds. Vol pp. 23-39, Wiley-Liss, Inc., New York, N.Y. 1994; Neben et al., Experimental Hematology 22:353-359, 1994; Cobblestone area forming cell assay, Ploemacher, R. E. In Culture of Hematopoietic Cells. R. I. Freshney, et al. eds. Vol pp. 1-21, Wiley-Liss, Inc., New York, N.Y. 1994; Long term bone marrow cultures in the presence of stromal cells, Spooncer, E., Dexter, M. and Allen, T. In Culture of Hematopoietic Cells. R. I. Freshney, et al. eds. Vol pp. 163-179, Wiley-Liss, Inc., New York, N.Y. 1994; Long term culture initiating cell assay, Sutherland, H. J. In Culture of Hematopoietic Cells. R. I. Freshney, et al. eds. Vol pp. 139-162, Wiley-Liss, Inc., New York, N.Y. 1994.

4.10.6 TISSUE GROWTH ACTIVITY

A polypeptide of the present invention also may be involved in bone, cartilage, tendon, ligament and/or nerve tissue growth or regeneration, as well as in wound healing and tissue repair and replacement, and in healing of burns, incisions and ulcers.

A polypeptide of the present invention which induces cartilage and/or bone growth in circumstances where bone is not normally formed, has application in the healing of bone fractures and cartilage damage or defects in humans and other animals. Compositions of a polypeptide, antibody, binding partner, or other modulator of the invention may have prophylactic use in closed as well as open fracture reduction and also in the improved fixation of artificial joints. De novo bone formation induced by an osteogenic agent contributes to the repair of congenital, trauma induced, or oncologic resection induced craniofacial defects, and also is useful in cosmetic plastic surgery.

A polypeptide of this invention may also be involved in attracting bone-forming cells, stimulating growth of bone-forming cells, or inducing differentiation of progenitors of bone-forming cells. Treatment of osteoporosis, osteoarthritis, bone degenerative disorders, or periodontal disease, such as through stimulation of bone and/or cartilage repair or by blocking inflammation or processes of tissue destruction (collagenase activity, osteoclast activity, etc.) mediated by inflammatory processes may also be possible using the composition of the invention.

Another category of tissue regeneration activity that may involve the polypeptide of the present invention is tendon/ligament formation. Induction of tendon/ligament-like tissue or other tissue formation in circumstances where such tissue is not normally formed, has application in the healing of tendon or ligament tears, deformities and other tendon or ligament defects in humans and other animals. Such a preparation employing a tendon/ligament-like tissue inducing protein may have prophylactic use in preventing damage to tendon or ligament tissue, as well as use in the improved fixation of tendon or ligament to bone or other tissues, and in repairing defects to tendon or ligament tissue. De novo tendon/ligament-like tissue formation induced by a composition of the present invention contributes to the repair of congenital, trauma induced, or other tendon or ligament defects of other origin, and is also useful in cosmetic plastic surgery for attachment or repair of tendons or ligaments. The compositions of the present invention may provide environment to attract tendon- or ligament-forming cells, stimulate growth of tendon- or ligament-forming cells, induce differentiation of progenitors of tendon- or ligament-forming cells, or induce growth of tendon/ligament cells or progenitors ex vivo for return in vivo to effect tissue repair. The compositions of the invention may also be useful in the treatment of tendinitis, carpal tunnel syndrome and other tendon or ligament defects. The compositions may also include an appropriate matrix and/or sequestering agent as a carrier as is well known in the art.

The compositions of the present invention may also be useful for proliferation of neural cells and for regeneration of nerve and brain tissue, i.e. for the treatment of central and peripheral nervous system diseases and neuropathies, as well as mechanical and traumatic disorders, which involve degeneration, death or trauma to neural cells or nerve tissue. More specifically, a composition may be used in the treatment of diseases of the peripheral nervous system, such as peripheral nerve injuries, peripheral neuropathy and localized neuropathies, and central nervous system diseases, such as Alzheimer's, Parkinson's disease, Huntington's disease, amyotrophic lateral sclerosis, and Shy-Drager syndrome. Further conditions which may be treated in accordance with the present invention include mechanical and traumatic disorders, such as spinal cord disorders, head trauma and cerebrovascular diseases such as stroke. Peripheral neuropathies

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resulting from chemotherapy or other medical therapies may also be treatable using a composition of the invention.

Compositions of the invention may also be useful to promote better or faster closure of non-healing wounds, including without limitation pressure ulcers, ulcers associated with vascular insufficiency, surgical and traumatic wounds, and the like.

Compositions of the present invention may also be involved in the generation or regeneration of other tissues, such as organs (including, for example, pancreas, liver, intestine, kidney, skin, endothelium), muscle (smooth, skeletal or cardiac) and vascular (including vascular endothelium) tissue, or for promoting the growth of cells comprising such tissues. Part of the desired effects may be by inhibition or modulation of fibrotic scarring may allow normal tissue to regenerate. A polypeptide of the present invention may also exhibit angiogenic activity.

A composition of the present invention may also be useful for gut protection or regeneration and treatment of lung or liver fibrosis, reperfusion injury in various tissues, and conditions resulting from systemic cytokine damage.

A composition of the present invention may also be useful for promoting or inhibiting differentiation of tissues described above from precursor tissues or cells; or for inhibiting the growth of tissues described above.

Therapeutic compositions of the invention can be used in the following:

Assays for tissue generation activity include, without limitation, those described in: International Patent Publication No. WO95/16035 (bone, cartilage, tendon); International Patent Publication No. WO95/05846 (nerve, neuronal); International Patent Publication No. WO91/07491 (skin, endothelium).

Assays for wound healing activity include, without limitation, those described in: Winter, Epidermal Wound Healing, pps. 71-112 (Maibach, H. I. and Rovee, D. T., eds.), Year Book Medical Publishers, Inc., Chicago, as modified by Eaglstein and Mertz, J. Invest. Dermatol 71:382-84 (1978).

4.10.7 IMMUNE STIMULATING OR SUPPRESSING ACTIVITY

A polypeptide of the present invention may also exhibit immune stimulating or immune suppressing activity, including without limitation the activities for which assays are described herein. A polynucleotide of the invention can encode a polypeptide exhibiting such activities. A protein may be useful in the treatment of various immune deficiencies and disorders (including severe combined immunodeficiency (SCID)), e.g., in regulating (up or down) growth and proliferation of T and/or B lymphocytes, as well as effecting the cytolytic activity of NK cells and other cell populations. These immune deficiencies may be genetic or be caused by viral (e.g., HIV) as well as bacterial or fungal infections, or may result from autoimmune disorders. More specifically, infectious diseases causes by viral, bacterial, fungal or other infection may be treatable using a protein of the present invention, including infections by HIV, hepatitis viruses, herpes viruses, mycobacteria, Leishmania spp., malaria spp. and various fungal infections such as candidiasis. Of course, in this regard, proteins of the present invention may also be useful where a boost to the immune system generally may be desirable, i.e., in the treatment of cancer.

Autoimmune disorders which may be treated using a protein of the present invention include, for example, connective tissue disease, multiple sclerosis, systemic lupus erythematosus, rheumatoid arthritis, autoimmune pulmonary inflammation, Guillain-Barre syndrome, autoimmune thyroiditis, insulin dependent diabetes mellitis, myasthenia gravis, graft-versus-host disease and autoimmune inflammatory eye disease. Such a protein (or antagonists thereof, including antibodies) of the present invention may also to be useful in the treatment of allergic reactions and conditions (e.g., anaphylaxis, serum sickness, drug reactions, food allergies, insect venom allergies, mastocytosis, allergic rhinitis, hypersensitivity pneumonitis, urticaria, angioedema, eczema, atopic dermatitis, allergic contact dermatitis, erythema multiforme. Stevens-Johnson syndrome, allergic conjunctivitis, atopic keratoconjunctivitis, venereal keratoconjunctivitis, giant papillary conjunctivitis and contact allergies), such as asthma (particularly allergic asthma) or other respiratory problems. Other conditions, in which immune suppression is desired (including, for example, organ transplantation), may also be treatable using a protein (or antagonists thereof) of the present invention. The therapeutic effects of the polypeptides or antagonists thereof on allergic reactions can be evaluated by in vivo animals models such as the cumulative contact enhancement test (Lastborn et al., Toxicology 125: 59-66, 1998), skin prick test (Hoffmann et al., Allergy 54: 446-54, 1999), guinea pig skin sensitization test (Vohr et al., Arch. Toxocol. 73: 501-9), and murine local lymph node assay (Kimber et al., J. Toxicol. Environ. Health 53: 563-79).

Using the proteins of the invention it may also be possible to modulate immune responses, in a number of ways. Down regulation may be in the form of inhibiting or blocking an immune response already in progress or may involve preventing the induction of an immune response. The functions of activated T cells may be inhibited by suppressing T cell responses or by inducing specific tolerance in T cells, or both. Immunosuppression of T cell responses is generally an active, non-antigen-specific, process which requires continuous exposure of the T cells to the suppressive agent. Tolerance, which involves inducing non-responsiveness or anergy in T cells, is distinguishable from immunosuppression in that it is generally antigen-specific and persists after exposure to the tolerizing agent has ceased. Operationally, tolerance can be

demonstrated by the lack of a T cell response upon reexposure to specific antigen in the absence of the tolerizing agent.

Down regulating or preventing one or more antigen functions (including without limitation B lymphocyte antigen functions (such as, for example, B7)), e.g., preventing high level lymphokine synthesis by activated T cells, will be useful in situations of tissue, skin and organ transplantation and in graft-versus-host disease (GVHD). For example, blockage of T cell function should result in reduced tissue destruction in tissue transplantation. Typically, in tissue transplants, rejection of the transplant is initiated through its recognition as foreign by T cells, followed by an immune reaction that destroys the transplant. The administration of a therapeutic composition of the invention may prevent cytokine synthesis by immune cells, such as T cells, and thus acts as an immunosuppressant. Moreover, a lack of costimulation may also be sufficient to anergize the T cells, thereby inducing tolerance in a subject. Induction of long-term tolerance by B lymphocyte antigen-blocking reagents may avoid the necessity of repeated administration of these blocking reagents. To achieve sufficient immunosuppression or tolerance in a subject, it may also be necessary to block the function of a combination of B lymphocyte antigens.

The efficacy of particular therapeutic compositions in preventing organ transplant rejection or GVHD can be assessed using animal models that are predictive of efficacy in humans. Examples of appropriate systems which can be used include allogeneic cardiac grafts in rats and xenogeneic pancreatic islet cell grafts in mice, both of which have been used to examine the immunosuppressive effects of CTLA4Ig fusion proteins in vivo as described in Lenschow et al., Science 257:789-792 (1992) and Turka et al., Proc. Natl. Acad. Sci USA, 89:11102-11105 (1992). In addition, murine models of GVHD (see Paul ed., Fundamental Immunology, Raven Press, New York, 1989, pp. 846-847) can be used to determine the effect of therapeutic compositions of the invention on the development of that disease.

Blocking antigen function may also be therapeutically useful for treating autoimmune diseases. Many autoimmune disorders are the result of inappropriate activation of T cells that are reactive against self tissue and which promote the production of cytokines and autoantibodies involved in the pathology of the diseases. Preventing the activation of autoreactive T cells may reduce or eliminate disease symptoms. Administration of reagents which block stimulation of T cells can be used to inhibit T cell activation and prevent production of autoantibodies or T cell-derived cytokines which may be involved in the disease process. Additionally, blocking reagents may induce antigen-specific tolerance of autoreactive T cells which could lead to long-term relief from the disease. The efficacy of blocking reagents in preventing or alleviating autoimmune disorders can be determined using a number of well-characterized animal models of human autoimmune diseases. Examples include murine experimental autoimmune encephalitis,

systemic lupus erythmatosis in MRL/lpr/lpr mice or NZB hybrid mice, murine autoimmune collagen arthritis, diabetes mellitus in NOD mice and BB rats, and murine experimental myasthenia gravis (see Paul ed., Fundamental Immunology, Raven Press, New York, 1989, pp. 840-856).

Upregulation of an antigen function (e.g., a B lymphocyte antigen function), as a means of up regulating immune responses, may also be useful in therapy. Upregulation of immune responses may be in the form of enhancing an existing immune response or eliciting an initial immune response. For example, enhancing an immune response may be useful in cases of viral infection, including systemic viral diseases such as influenza, the common cold, and encephalitis.

Alternatively, anti-viral immune responses may be enhanced in an infected patient by removing T cells from the patient, costimulating the T cells in vitro with viral antigen-pulsed APCs either expressing a peptide of the present invention or together with a stimulatory form of a soluble peptide of the present invention and reintroducing the in vitro activated T cells into the patient. Another method of enhancing anti-viral immune responses would be to isolate infected cells from a patient, transfect them with a nucleic acid encoding a protein of the present invention as described herein such that the cells express all or a portion of the protein on their surface, and reintroduce the transfected cells into the patient. The infected cells would now be capable of delivering a costimulatory signal to, and thereby activate, T cells in vivo.

A polypeptide of the present invention may provide the necessary stimulation signal to T cells to induce a T cell mediated immune response against the transfected tumor cells. In addition, tumor cells which lack MHC class I or MHC class II molecules, or which fail to reexpress sufficient mounts of MHC class I or MHC class II molecules, can be transfected with nucleic acid encoding all or a portion of (e.g., a cytoplasmic-domain truncated portion) of an MHC class I alpha chain protein and β₂ microglobulin protein or an MHC class II alpha chain protein and an MHC class II beta chain protein to thereby express MHC class I or MHC class II proteins on the cell surface. Expression of the appropriate class I or class II MHC in conjunction with a peptide having the activity of a B lymphocyte antigen (e.g., B7-1, B7-2, B7-3) induces a T cell mediated immune response against the transfected tumor cell. Optionally, a gene encoding an antisense construct which blocks expression of an MHC class II associated protein, such as the invariant chain, can also be cotransfected with a DNA encoding a peptide having the activity of a B lymphocyte antigen to promote presentation of tumor associated antigens and induce tumor specific immunity. Thus, the induction of a T cell mediated immune response in a human subject may be sufficient to overcome tumor-specific tolerance in the subject.

The activity of a protein of the invention may, among other means, be measured by the following methods:

Suitable assays for thymocyte or splenocyte cytotoxicity include, without limitation, those described in: Current Protocols in Immunology, Ed by J. E. Coligan, A. M. Kruisbeek, D. H. Margulies, E. M. Shevach, W. Strober, Pub. Greene Publishing Associates and Wiley-Interscience (Chapter 3, In Vitro assays for Mouse Lymphocyte Function 3.1-3.19; Chapter 7, Immunologic studies in Humans); Herrmann et al., Proc. Natl. Acad. Sci. USA 78:2488-2492, 1981; Herrmann et al., J. Immunol. 128:1968-1974, 1982; Handa et al., J. Immunol. 135:1564-1572, 1985; Takai et al., I. Immunol. 137:3494-3500, 1986; Takai et al., J. Immunol. 140:508-512, 1988; Bowman et al., J. Virology 61:1992-1998; Bertagnolli et al., Cellular Immunology 133:327-341, 1991; Brown et al., J. Immunol. 153:3079-3092, 1994.

Assays for T-cell-dependent immunoglobulin responses and isotype switching (which will identify, among others, proteins that modulate T-cell dependent antibody responses and that affect Th1/Th2 profiles) include, without limitation, those described in: Maliszewski, J. Immunol. 144:3028-3033, 1990; and Assays for B cell function: In vitro antibody production, Mond, J. J. and Brunswick, M. In Current Protocols in Immunology. J. E. e.a. Coligan eds. Vol 1 pp. 3.8.1-3.8.16, John Wiley and Sons, Toronto. 1994.

Mixed lymphocyte reaction (MLR) assays (which will identify, among others, proteins that generate predominantly Th1 and CTL responses) include, without limitation, those described in: Current Protocols in Immunology, Ed by J. E. Coligan, A. M. Kruisbeek, D. H. Margulies, E. M. Shevach, W. Strober, Pub. Greene Publishing Associates and Wiley-Interscience (Chapter 3, In Vitro assays for Mouse Lymphocyte Function 3.1-3.19; Chapter 7, Immunologic studies in Humans); Takai et al., J. Immunol. 137:3494-3500, 1986; Takai et al., J. Immunol. 140:508-512, 1988; Bertagnolli et al., J. Immunol. 149:3778-3783, 1992.

Dendritic cell-dependent assays (which will identify, among others, proteins expressed by dendritic cells that activate naive T-cells) include, without limitation, those described in: Guery et al., J. Immunol. 134:536-544, 1995; Inaba et al., Journal of Experimental Medicine 173:549-559, 1991; Macatonia et al., Journal of Immunology 154:5071-5079, 1995; Porgador et al., Journal of Experimental Medicine 182:255-260, 1995; Nair et al., Journal of Virology 67:4062-4069, 1993; Huang et al., Science 264:961-965, 1994; Macatonia et al., Journal of Experimental Medicine 169:1255-1264, 1989; Bhardwaj et al., Journal of Clinical Investigation 94:797-807, 1994; and Inaba et al., Journal of Experimental Medicine 172:631-640, 1990.

Assays for lymphocyte survival/apoptosis (which will identify, among others, proteins that prevent apoptosis after superantigen induction and proteins that regulate lymphocyte homeostasis) include, without limitation, those described in: Darzynkiewicz et al., Cytometry

13:795-808, 1992; Gorczyca et al., Leukemia 7:659-670, 1993; Gorczyca et al., Cancer Research 53:1945-1951, 1993; Itoh et al., Cell 66:233-243, 1991; Zacharchuk, Journal of Immunology 145:4037-4045, 1990; Zamai et al., Cytometry 14:891-897, 1993; Gorczyca et al., International Journal of Oncology 1:639-648, 1992.

Assays for proteins that influence early steps of T-cell commitment and development include, without limitation, those described in: Antica et al., Blood 84:111-117, 1994; Fine et al., Cellular Immunology 155:111-122, 1994; Galy et al., Blood 85:2770-2778, 1995; Toki et al., Proc. Nat. Acad Sci. USA 88:7548-7551, 1991.

4.10.8 ACTIVIN/INHIBIN ACTIVITY

A polypeptide of the present invention may also exhibit activin- or inhibin-related activities. A polynucleotide of the invention may encode a polypeptide exhibiting such characteristics. Inhibins are characterized by their ability to inhibit the release of follicle stimulating hormone (FSH), while activins and are characterized by their ability to stimulate the release of follicle stimulating hormone (FSH). Thus, a polypeptide of the present invention, alone or in heterodimers with a member of the inhibin family, may be useful as a contraceptive based on the ability of inhibins to decrease fertility in female mammals and decrease spermatogenesis in male mammals. Administration of sufficient amounts of other inhibins can induce infertility in these mammals. Alternatively, the polypeptide of the invention, as a homodimer or as a heterodimer with other protein subunits of the inhibin group, may be useful as a fertility inducing therapeutic, based upon the ability of activin molecules in stimulating FSH release from cells of the anterior pituitary. See, for example, U.S. Pat. No. 4,798,885. A polypeptide of the invention may also be useful for advancement of the onset of fertility in sexually immature mammals, so as to increase the lifetime reproductive performance of domestic animals such as, but not limited to, cows, sheep and pigs.

The activity of a polypeptide of the invention may, among other means, be measured by the following methods.

Assays for activin/inhibin activity include, without limitation, those described in: Vale et al., Endocrinology 91:562-572, 1972; Ling et al., Nature 321:779-782, 1986; Vale et al., Nature 321:776-779, 1986; Mason et al., Nature 318:659-663, 1985; Forage et al., Proc. Natl. Acad. Sci. USA 83:3091-3095, 1986.

4.10.9 CHEMOTACTIC/CHEMOKINETIC ACTIVITY

A polypeptide of the present invention may be involved in chemotactic or chemokinetic activity for mammalian cells, including, for example, monocytes, fibroblasts, neutrophils,

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T-cells, mast cells, eosinophils, epithelial and/or endothelial cells. A polynucleotide of the invention can encode a polypeptide exhibiting such attributes. Chemotactic and chemokinetic receptor activation can be used to mobilize or attract a desired cell population to a desired site of action. Chemotactic or chemokinetic compositions (e.g. proteins, antibodies, binding partners, or modulators of the invention) provide particular advantages in treatment of wounds and other trauma to tissues, as well as in treatment of localized infections. For example, attraction of lymphocytes, monocytes or neutrophils to tumors or sites of infection may result in improved immune responses against the tumor or infecting agent.

A protein or peptide has chemotactic activity for a particular cell population if it can stimulate, directly or indirectly, the directed orientation or movement of such cell population. Preferably, the protein or peptide has the ability to directly stimulate directed movement of cells. Whether a particular protein has chemotactic activity for a population of cells can be readily determined by employing such protein or peptide in any known assay for cell chemotaxis.

Therapeutic compositions of the invention can be used in the following:

Assays for chemotactic activity (which will identify proteins that induce or prevent chemotaxis) consist of assays that measure the ability of a protein to induce the migration of cells across a membrane as well as the ability of a protein to induce the adhesion of one cell population to another cell population. Suitable assays for movement and adhesion include, without limitation, those described in: Current Protocols in Immunology, Ed by J. E. Coligan, A. M. Kruisbeek, D. H. Marguiles, E. M. Shevach, W. Strober, Pub. Greene Publishing Associates and Wiley-Interscience (Chapter 6.12, Measurement of alpha and beta Chemokines 6.12.1-6.12.28; Taub et al. J. Clin. Invest. 95:1370-1376, 1995; Lind et al. APMIS 103:140-146, 1995; Muller et al Eur. J. Immunol. 25:1744-1748; Gruber et al. J. of Immunol. 152:5860-5867, 1994; Johnston et al. J. of Immunol. 153:1762-1768, 1994.

4.10.10 HEMOSTATIC AND THROMBOLYTIC ACTIVITY

A polypeptide of the invention may also be involved in hemostasis or thrombolysis or thrombosis. A polynucleotide of the invention can encode a polypeptide exhibiting such attributes. Compositions may be useful in treatment of various coagulation disorders (including hereditary disorders, such as hemophilias) or to enhance coagulation and other hemostatic events in treating wounds resulting from trauma, surgery or other causes. A composition of the invention may also be useful for dissolving or inhibiting formation of thromboses and for treatment and prevention of conditions resulting therefrom (such as, for example, infarction of cardiac and central nervous system vessels (e.g., stroke).

Therapeutic compositions of the invention can be used in the following:

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Assay for hemostatic and thrombolytic activity include, without limitation, those described in: Linet et al., J. Clin. Pharmacol. 26:131-140, 1986; Burdick et al., Thrombosis Res. 45:413-419, 1987; Humphrey et al., Fibrinolysis 5:71-79 (1991); Schaub, Prostaglandins 35:467-474, 1988.

4.10.11 **CANCER DIAGNOSIS AND THERAPY**

Polypeptides of the invention may be involved in cancer cell generation, proliferation or metastasis. Detection of the presence or amount of polynucleotides or polypeptides of the invention may be useful for the diagnosis and/or prognosis of one or more types of cancer. For example, the presence or increased expression of a polynucleotide/polypeptide of the invention may indicate a hereditary risk of cancer, a precancerous condition, or an ongoing malignancy. Conversely, a defect in the gene or absence of the polypeptide may be associated with a cancer condition. Identification of single nucleotide polymorphisms associated with cancer or a predisposition to cancer may also be useful for diagnosis or prognosis.

Cancer treatments promote tumor regression by inhibiting tumor cell proliferation, inhibiting angiogenesis (growth of new blood vessels that is necessary to support tumor growth) and/or prohibiting metastasis by reducing tumor cell motility or invasiveness. Therapeutic compositions of the invention may be effective in adult and pediatric oncology including in solid phase tumors/malignancies, locally advanced tumors, human soft tissue sarcomas, metastatic cancer, including lymphatic metastases, blood cell malignancies including multiple myeloma, acute and chronic leukemias, and lymphomas, head and neck cancers including mouth cancer, larynx cancer and thyroid cancer, lung cancers including small cell carcinoma and non-small cell cancers, breast cancers including small cell carcinoma and ductal carcinoma, gastrointestinal cancers including esophageal cancer, stomach cancer, colon cancer, colorectal cancer and polyps associated with colorectal neoplasia, pancreatic cancers, liver cancer, urologic cancers including bladder cancer and prostate cancer, malignancies of the female genital tract including ovarian carcinoma, uterine (including endometrial) cancers, and solid tumor in the ovarian follicle, kidney cancers including renal cell carcinoma, brain cancers including intrinsic brain tumors, neuroblastoma, astrocytic brain tumors, gliomas, metastatic tumor cell invasion in the central nervous system, bone cancers including osteomas, skin cancers including malignant melanoma, tumor progression of human skin keratinocytes, squamous cell carcinoma, basal cell carcinoma, hemangiopericytoma and Karposi's sarcoma.

Polypeptides, polynucleotides, or modulators of polypeptides of the invention (including inhibitors and stimulators of the biological activity of the polypeptide of the invention) may be administered to treat cancer. Therapeutic compositions can be administered in therapeutically

effective dosages alone or in combination with adjuvant cancer therapy such as surgery, chemotherapy, radiotherapy, thermotherapy, and laser therapy, and may provide a beneficial effect, e.g. reducing tumor size, slowing rate of tumor growth, inhibiting metastasis, or otherwise improving overall clinical condition, without necessarily eradicating the cancer.

The composition can also be administered in therapeutically effective amounts as a portion of an anti-cancer cocktail. An anti-cancer cocktail is a mixture of the polypeptide or modulator of the invention with one or more anti-cancer drugs in addition to a pharmaceutically acceptable carrier for delivery. The use of anti-cancer cocktails as a cancer treatment is routine. Anti-cancer drugs that are well known in the art and can be used as a treatment in combination with the polypeptide or modulator of the invention include: Actinomycin D, Aminoglutethimide, Asparaginase, Bleomycin, Busulfan, Carboplatin, Carmustine, Chlorambucil, Cisplatin (cis-DDP), Cyclophosphamide, Cytarabine HCl (Cytosine arabinoside), Dacarbazine, Dactinomycin, Daunorubicin HCl, Doxorubicin HCl, Estramustine phosphate sodium, Etoposide (V16-213), Floxuridine, 5-Fluorouracil (5-Fu), Flutamide, Hydroxyurea (hydroxycarbamide), Ifosfamide. Interferon Alpha-2a, Interferon Alpha-2b, Leuprolide acetate (LHRH-releasing factor analog), Lomustine, Mechlorethamine HCl (nitrogen mustard), Melphalan, Mercaptopurine, Mesna, Methotrexate (MTX), Mitomycin, Mitoxantrone HCl, Octreotide, Plicamycin, Procarbazine HCl, Streptozocin, Tamoxifen citrate, Thioguanine, Thiotepa, Vinblastine sulfate, Vincristine sulfate, Amsacrine, Azacitidine, Hexamethylmelamine, Interleukin-2, Mitoguazone, Pentostatin, Semustine, Teniposide, and Vindesine sulfate.

In addition, therapeutic compositions of the invention may be used for prophylactic treatment of cancer. There are hereditary conditions and/or environmental situations (e.g. exposure to carcinogens) known in the art that predispose an individual to developing cancers. Under these circumstances, it may be beneficial to treat these individuals with therapeutically effective doses of the polypeptide of the invention to reduce the risk of developing cancers.

In vitro models can be used to determine the effective doses of the polypeptide of the invention as a potential cancer treatment. These in vitro models include proliferation assays of cultured tumor cells, growth of cultured tumor cells in soft agar (see Freshney, (1987) Culture of Animal Cells: A Manual of Basic Technique, Wily-Liss, New York, NY Ch 18 and Ch 21), tumor systems in nude mice as described in Giovanella et al., J. Natl. Can. Inst., 52: 921-30 (1974), mobility and invasive potential of tumor cells in Boyden Chamber assays as described in Pilkington et al., Anticancer Res., 17: 4107-9 (1997), and angiogenesis assays such as induction of vascularization of the chick chorioallantoic membrane or induction of vascular endothelial cell migration as described in Ribatta et al., Intl. J. Dev. Biol., 40: 1189-97 (1999) and Li et al., Clin. Exp. Metastasis, 17:423-9 (1999), respectively. Suitable tumor cells lines are available,

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e.g. from American Type Tissue Culture Collection catalogs.

4.10.12 RECEPTOR/LIGAND ACTIVITY

A polypeptide of the present invention may also demonstrate activity as receptor, receptor ligand or inhibitor or agonist of receptor/ligand interactions. A polynucleotide of the invention can encode a polypeptide exhibiting such characteristics. Examples of such receptors and ligands include, without limitation, cytokine receptors and their ligands, receptor kinases and their ligands, receptor phosphatases and their ligands, receptors involved in cell-cell interactions and their ligands (including without limitation, cellular adhesion molecules (such as selectins, integrins and their ligands) and receptor/ligand pairs involved in antigen presentation, antigen recognition and development of cellular and humoral immune responses. Receptors and ligands are also useful for screening of potential peptide or small molecule inhibitors of the relevant receptor/ligand interaction. A protein of the present invention (including, without limitation, fragments of receptors and ligands) may themselves be useful as inhibitors of receptor/ligand interactions.

The activity of a polypeptide of the invention may, among other means, be measured by the following methods:

Suitable assays for receptor-ligand activity include without limitation those described in: Current Protocols in Immunology, Ed by J. E. Coligan, A. M. Kruisbeek, D. H. Margulies, E. M. Shevach, W. Strober, Pub. Greene Publishing Associates and Wiley- Interscience (Chapter 7.28, Measurement of Cellular Adhesion under static conditions 7.28.1-7.28.22), Takai et al., Proc. Natl. Acad. Sci. USA 84:6864-6868, 1987; Bierer et al., J. Exp. Med. 168:1145-1156, 1988; Rosenstein et al., J. Exp. Med. 169:149-160 1989; Stoltenborg et al., J. Immunol. Methods 175:59-68, 1994; Stitt et al., Cell 80:661-670, 1995.

By way of example, the polypeptides of the invention may be used as a receptor for a ligand(s) thereby transmitting the biological activity of that ligand(s). Ligands may be identified through binding assays, affinity chromatography, dihybrid screening assays, BIAcore assays, gel overlay assays, or other methods known in the art.

Studies characterizing drugs or proteins as agonist or antagonist or partial agonists or a partial antagonist require the use of other proteins as competing ligands. The polypeptides of the present invention or ligand(s) thereof may be labeled by being coupled to radioisotopes, colorimetric molecules or a toxin molecules by conventional methods. ("Guide to Protein Purification" Murray P. Deutscher (ed) Methods in Enzymology Vol. 182 (1990) Academic Press, Inc. San Diego). Examples of radioisotopes include, but are not limited to, tritium and carbon-14. Examples of colorimetric molecules include, but are not limited to, fluorescent

molecules such as fluorescamine, or rhodamine or other colorimetric molecules. Examples of toxins include, but are not limited, to ricin.

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4.10.13 DRUG SCREENING

This invention is particularly useful for screening chemical compounds by using the novel polypeptides or binding fragments thereof in any of a variety of drug screening techniques. The polypeptides or fragments employed in such a test may either be free in solution, affixed to a solid support, borne on a cell surface or located intracellularly. One method of drug screening utilizes eukaryotic or prokaryotic host cells which are stably transformed with recombinant nucleic acids expressing the polypeptide or a fragment thereof. Drugs are screened against such transformed cells in competitive binding assays. Such cells, either in viable or fixed form, can be used for standard binding assays. One may measure, for example, the formation of complexes between polypeptides of the invention or fragments and the agent being tested or examine the diminution in complex formation between the novel polypeptides and an appropriate cell line, which are well known in the art.

Sources for test compounds that may be screened for ability to bind to or modulate (i.e., increase or decrease) the activity of polypeptides of the invention include (1) inorganic and organic chemical libraries, (2) natural product libraries, and (3) combinatorial libraries comprised of either random or mimetic peptides, oligonucleotides or organic molecules.

Chemical libraries may be readily synthesized or purchased from a number of commercial sources, and may include structural analogs of known compounds or compounds that are identified as "hits" or "leads" via natural product screening.

The sources of natural product libraries are microorganisms (including bacteria and fungi), animals, plants or other vegetation, or marine organisms, and libraries of mixtures for screening may be created by: (1) fermentation and extraction of broths from soil, plant or marine microorganisms or (2) extraction of the organisms themselves. Natural product libraries include polyketides, non-ribosomal peptides, and (non-naturally occurring) variants thereof. For a review, see *Science* 282:63-68 (1998).

Combinatorial libraries are composed of large numbers of peptides, oligonucleotides or organic compounds and can be readily prepared by traditional automated synthesis methods, PCR, cloning or proprietary synthetic methods. Of particular interest are peptide and oligonucleotide combinatorial libraries. Still other libraries of interest include peptide, protein, peptidomimetic, multiparallel synthetic collection, recombinatorial, and polypeptide libraries. For a review of combinatorial chemistry and libraries created therefrom, see Myers, Curr. Opin. Biotechnol. 8:701-707 (1997). For reviews and examples of peptidomimetic libraries, see

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Al-Obeidi et al., Mol. Biotechnol, 9(3):205-23 (1998); Hruby et al., Curr Opin Chem Biol, 1(1):114-19 (1997); Dorner et al., Bioorg Med Chem, 4(5):709-15 (1996) (alkylated dipeptides).

Identification of modulators through use of the various libraries described herein permits modification of the candidate "hit" (or "lead") to optimize the capacity of the "hit" to bind a polypeptide of the invention. The molecules identified in the binding assay are then tested for antagonist or agonist activity in in vivo tissue culture or animal models that are well known in the art. In brief, the molecules are titrated into a plurality of cell cultures or animals and then tested for either cell/animal death or prolonged survival of the animal/cells.

The binding molecules thus identified may be complexed with toxins, e.g., ricin or cholera, or with other compounds that are toxic to cells such as radioisotopes. The toxin-binding molecule complex is then targeted to a tumor or other cell by the specificity of the binding molecule for a polypeptide of the invention. Alternatively, the binding molecules may be complexed with imaging agents for targeting and imaging purposes.

4.10.14 ASSAY FOR RECEPTOR ACTIVITY

The invention also provides methods to detect specific binding of a polypeptide e.g. a ligand or a receptor. The art provides numerous assays particularly useful for identifying previously unknown binding partners for receptor polypeptides of the invention. For example, expression cloning using mammalian or bacterial cells, or dihybrid screening assays can be used to identify polynucleotides encoding binding partners. As another example, affinity chromatography with the appropriate immobilized polypeptide of the invention can be used to isolate polypeptides that recognize and bind polypeptides of the invention. There are a number of different libraries used for the identification of compounds, and in particular small molecules, that modulate (i.e., increase or decrease) biological activity of a polypeptide of the invention. Ligands for receptor polypeptides of the invention can also be identified by adding exogenous ligands, or cocktails of ligands to two cells populations that are genetically identical except for the expression of the receptor of the invention: one cell population expresses the receptor of the invention whereas the other does not. The response of the two cell populations to the addition of ligands(s) are then compared. Alternatively, an expression library can be co-expressed with the polypeptide of the invention in cells and assayed for an autocrine response to identify potential ligand(s). As still another example, BIAcore assays, gel overlay assays, or other methods known in the art can be used to identify binding partner polypeptides, including, (1) organic and inorganic chemical libraries, (2) natural product libraries, and (3) combinatorial libraries comprised of random peptides, oligonucleotides or organic molecules.

The role of downstream intracellular signaling molecules in the signaling cascade of the polypeptide of the invention can be determined. For example, a chimeric protein in which the cytoplasmic domain of the polypeptide of the invention is fused to the extracellular portion of a protein, whose ligand has been identified, is produced in a host cell. The cell is then incubated with the ligand specific for the extracellular portion of the chimeric protein, thereby activating the chimeric receptor. Known downstream proteins involved in intracellular signaling can then be assayed for expected modifications i.e. phosphorylation. Other methods known to those in the art can also be used to identify signaling molecules involved in receptor activity.

4.10.15 ANTI-INFLAMMATORY ACTIVITY

Compositions of the present invention may also exhibit anti-inflammatory activity. The anti-inflammatory activity may be achieved by providing a stimulus to cells involved in the inflammatory response, by inhibiting or promoting cell-cell interactions (such as, for example, cell adhesion), by inhibiting or promoting chemotaxis of cells involved in the inflammatory process, inhibiting or promoting cell extravasation, or by stimulating or suppressing production of other factors which more directly inhibit or promote an inflammatory response. Compositions with such activities can be used to treat inflammatory conditions including chronic or acute conditions), including without limitation intimation associated with infection (such as septic shock, sepsis or systemic inflammatory response syndrome (SIRS)), ischemia-reperfusion injury, endotoxin lethality, arthritis, complement-mediated hyperacute rejection, nephritis, cytokine or chemokine-induced lung injury, inflammatory bowel disease, Crohn's disease or resulting from over production of cytokines such as TNF or IL-1. Compositions of the invention may also be useful to treat anaphylaxis and hypersensitivity to an antigenic substance or material. Compositions of this invention may be utilized to prevent or treat conditions such as, but not limited to, sepsis, acute pancreatitis, endotoxin shock, cytokine induced shock, rheumatoid arthritis, chronic inflammatory arthritis, pancreatic cell damage from diabetes mellitus type 1, graft versus host disease, inflammatory bowel disease, inflamation associated with pulmonary disease, other autoimmune disease or inflammatory disease, an antiproliferative agent such as for acute or chronic mylegenous leukemia or in the prevention of premature labor secondary to intrauterine infections.

4.10.16 LEUKEMIAS

Leukemias and related disorders may be treated or prevented by administration of a therapeutic that promotes or inhibits function of the polynucleotides and/or polypeptides of the invention. Such leukemias and related disorders include but are not limited to acute leukemia,

acute lymphocytic leukemia, acute myelocytic leukemia, myeloblastic, promyelocytic, myelomonocytic, monocytic, erythroleukemia, chronic leukemia, chronic myelocytic (granulocytic) leukemia and chronic lymphocytic leukemia (for a review of such disorders, see Fishman et al., 1985, Medicine, 2d Ed., J.B. Lippincott Co., Philadelphia).

4.10.17 NERVOUS SYSTEM DISORDERS

Nervous system disorders, involving cell types which can be tested for efficacy of intervention with compounds that modulate the activity of the polynucleotides and/or polypeptides of the invention, and which can be treated upon thus observing an indication of therapeutic utility, include but are not limited to nervous system injuries, and diseases or disorders which result in either a disconnection of axons, a diminution or degeneration of neurons, or demyelination. Nervous system lesions which may be treated in a patient (including human and non-human mammalian patients) according to the invention include but are not limited to the following lesions of either the central (including spinal cord, brain) or peripheral nervous systems:

- (i) traumatic lesions, including lesions caused by physical injury or associated with surgery, for example, lesions which sever a portion of the nervous system, or compression injuries;
- (ii) ischemic lesions, in which a lack of oxygen in a portion of the nervous system
 results in neuronal injury or death, including cerebral infarction or ischemia, or spinal cord
 infarction or ischemia;
- (iii) infectious lesions, in which a portion of the nervous system is destroyed or injured as a result of infection, for example, by an abscess or associated with infection by human immunodeficiency virus, herpes zoster, or herpes simplex virus or with Lyme disease, tuberculosis, syphilis;
- (iv) degenerative lesions, in which a portion of the nervous system is destroyed or injured as a result of a degenerative process including but not limited to degeneration associated with Parkinson's disease, Alzheimer's disease, Huntington's chorea, or amyotrophic lateral sclerosis;
- (v) lesions associated with nutritional diseases or disorders, in which a portion of the nervous system is destroyed or injured by a nutritional disorder or disorder of metabolism including but not limited to, vitamin B12 deficiency, folic acid deficiency, Wernicke disease, tobacco-alcohol amblyopia, Marchiafava-Bignami disease (primary degeneration of the corpus callosum), and alcoholic cerebellar degeneration;

- (vi) neurological lesions associated with systemic diseases including but not limited to diabetes (diabetic neuropathy, Bell's palsy), systemic lupus erythematosus, carcinoma, or sarcoidosis;
- (vii) lesions caused by toxic substances including alcohol, lead, or particular neurotoxins; and
- (viii) demyelinated lesions in which a portion of the nervous system is destroyed or injured by a demyelinating disease including but not limited to multiple sclerosis, human immunodeficiency virus-associated myelopathy, transverse myelopathy or various etiologies, progressive multifocal leukoencephalopathy, and central pontine myelinolysis.

Therapeutics which are useful according to the invention for treatment of a nervous system disorder may be selected by testing for biological activity in promoting the survival or differentiation of neurons. For example, and not by way of limitation, therapeutics which elicit any of the following effects may be useful according to the invention:

- (i) increased survival time of neurons in culture;
- (ii) increased sprouting of neurons in culture or in vivo;
- (iii) increased production of a neuron-associated molecule in culture or *in vivo*, *e.g.*, choline acetyltransferase or acetylcholinesterase with respect to motor neurons; or
 - .(iv) decreased symptoms of neuron dysfunction in vivo.

Such effects may be measured by any method known in the art. In preferred, non-limiting embodiments, increased survival of neurons may be measured by the method set forth in Arakawa et al. (1990, J. Neurosci. 10:3507-3515); increased sprouting of neurons may be detected by methods set forth in Pestronk et al. (1980, Exp. Neurol. 70:65-82) or Brown et al. (1981, Ann. Rev. Neurosci. 4:17-42); increased production of neuron-associated molecules may be measured by bioassay, enzymatic assay, antibody binding, Northern blot assay, etc., depending on the molecule to be measured; and motor neuron dysfunction may be measured by assessing the physical manifestation of motor neuron disorder, e.g., weakness, motor neuron conduction velocity, or functional disability.

In specific embodiments, motor neuron disorders that may be treated according to the invention include but are not limited to disorders such as infarction, infection, exposure to toxin, trauma, surgical damage, degenerative disease or malignancy that may affect motor neurons as well as other components of the nervous system, as well as disorders that selectively affect neurons such as amyotrophic lateral sclerosis, and including but not limited to progressive spinal muscular atrophy, progressive bulbar palsy, primary lateral sclerosis, infantile and juvenile muscular atrophy, progressive bulbar paralysis of childhood (Fazio-Londe syndrome),

poliomyelitis and the post polio syndrome, and Hereditary Motorsensory Neuropathy (Charcot-Marie-Tooth Disease).

4.10.18 OTHER ACTIVITIES

A polypeptide of the invention may also exhibit one or more of the following additional activities or effects: inhibiting the growth, infection or function of, or killing, infectious agents, including, without limitation, bacteria, viruses, fungi and other parasites; effecting (suppressing or enhancing) bodily characteristics, including, without limitation, height, weight, hair color, eye color, skin, fat to lean ratio or other tissue pigmentation, or organ or body part size or shape (such as, for example, breast augmentation or diminution, change in bone form or shape); effecting biorhythms or circadian cycles or rhythms; effecting the fertility of male or female subjects; effecting the metabolism, catabolism, anabolism, processing, utilization, storage or elimination of dietary fat, lipid, protein, carbohydrate, vitamins, minerals, co-factors or other nutritional factors or component(s); effecting behavioral characteristics, including, without limitation, appetite, libido, stress, cognition (including cognitive disorders), depression (including depressive disorders) and violent behaviors; providing analgesic effects or other pain reducing effects; promoting differentiation and growth of embryonic stem cells in lineages other than hematopoietic lineages; hormonal or endocrine activity; in the case of enzymes, correcting deficiencies of the enzyme and treating deficiency-related diseases; treatment of hyperproliferative disorders (such as, for example, psoriasis); immunoglobulin-like activity (such as, for example, the ability to bind antigens or complement); and the ability to act as an antigen in a vaccine composition to raise an immune response against such protein or another material or entity which is cross-reactive with such protein.

4.10.19 IDENTIFICATION OF POLYMORPHISMS

The demonstration of polymorphisms makes possible the identification of such polymorphisms in human subjects and the pharmacogenetic use of this information for diagnosis and treatment. Such polymorphisms may be associated with, e.g., differential predisposition or susceptibility to various disease states (such as disorders involving inflammation or immune response) or a differential response to drug administration, and this genetic information can be used to tailor preventive or therapeutic treatment appropriately. For example, the existence of a polymorphism associated with a predisposition to inflammation or autoimmune disease makes possible the diagnosis of this condition in humans by identifying the presence of the polymorphism.

Polymorphisms can be identified in a variety of ways known in the art which all generally involve obtaining a sample from a patient, analyzing DNA from the sample, optionally involving isolation or amplification of the DNA, and identifying the presence of the polymorphism in the DNA. For example, PCR may be used to amplify an appropriate fragment of genomic DNA which may then be sequenced. Alternatively, the DNA may be subjected to allele-specific oligonucleotide hybridization (in which appropriate oligonucleotides are hybridized to the DNA under conditions permitting detection of a single base mismatch) or to a single nucleotide extension assay (in which an oligonucleotide that hybridizes immediately adjacent to the position of the polymorphism is extended with one or more labeled nucleotides). In addition, traditional restriction fragment length polymorphism analysis (using restriction enzymes that provide differential digestion of the genomic DNA depending on the presence or absence of the polymorphism) may be performed. Arrays with nucleotide sequences of the present invention can be used to detect polymorphisms. The array can comprise modified nucleotide sequences of the present invention in order to detect the nucleotide sequences of the present invention. In the alternative, any one of the nucleotide sequences of the present invention can be placed on the array to detect changes from those sequences.

Alternatively a polymorphism resulting in a change in the amino acid sequence could also be detected by detecting a corresponding change in amino acid sequence of the protein, e.g., by an antibody specific to the variant sequence.

4.10.20 ARTHRITIS AND INFLAMMATION

The immunosuppressive effects of the compositions of the invention against rheumatoid arthritis is determined in an experimental animal model system. The experimental model system is adjuvant induced arthritis in rats, and the protocol is described by J. Holoshitz, et at., 1983, Science, 219:56, or by B. Waksman et al., 1963, Int. Arch. Allergy Appl. Immunol., 23:129. Induction of the disease can be caused by a single injection, generally intradermally, of a suspension of killed Mycobacterium tuberculosis in complete Freund's adjuvant (CFA). The route of injection can vary, but rats may be injected at the base of the tail with an adjuvant mixture. The polypeptide is administered in phosphate buffered solution (PBS) at a dose of about 1-5 mg/kg. The control consists of administering PBS only.

The procedure for testing the effects of the test compound would consist of intradermally injecting killed Mycobacterium tuberculosis in CFA followed by immediately administering the test compound and subsequent treatment every other day until day 24. At 14, 15, 18, 20, 22, and 24 days after injection of Mycobacterium CFA, an overall arthritis score may be obtained as described by J. Holoskitz above. An analysis of the data would reveal that the test compound

would have a dramatic affect on the swelling of the joints as measured by a decrease of the arthritis score.

4.11 THERAPEUTIC METHODS

The compositions (including polypeptide fragments, analogs, variants and antibodies or other binding partners or modulators including antisense polynucleotides) of the invention have numerous applications in a variety of therapeutic methods. Examples of therapeutic applications include, but are not limited to, those exemplified herein.

4.11.1 EXAMPLE

One embodiment of the invention is the administration of an effective amount of the polypeptides or other composition of the invention to individuals affected by a disease or disorder that can be modulated by regulating the peptides of the invention. While the mode of administration is not particularly important, parenteral administration is preferred. An exemplary mode of administration is to deliver an intravenous bolus. The dosage of the polypeptides or other composition of the invention will normally be determined by the prescribing physician. It is to be expected that the dosage will vary according to the age, weight, condition and response of the individual patient. Typically, the amount of polypeptide administered per dose will be in the range of about 0.01µg/kg to 100 mg/kg of body weight, with the preferred dose being about 0.1µg/kg to 10 mg/kg of patient body weight. For parenteral administration, polypeptides of the invention will be formulated in an injectable form combined with a pharmaceutically acceptable parenteral vehicle. Such vehicles are well known in the art and examples include water, saline, Ringer's solution, dextrose solution, and solutions consisting of small amounts of the human serum albumin. The vehicle may contain minor amounts of additives that maintain the isotonicity and stability of the polypeptide or other active ingredient. The preparation of such solutions is within the skill of the art.

4.12 PHARMACEUTICAL FORMULATIONS AND ROUTES OF ADMINISTRATION

A protein or other composition of the present invention (from whatever source derived, including without limitation from recombinant and non-recombinant sources and including antibodies and other binding partners of the polypeptides of the invention) may be administered to a patient in need, by itself, or in pharmaceutical compositions where it is mixed with suitable carriers or excipient(s) at doses to treat or ameliorate a variety of disorders. Such a composition may optionally contain (in addition to protein or other active ingredient and a carrier) diluents,

fillers, salts, buffers, stabilizers, solubilizers, and other materials well known in the art. The term "pharmaceutically acceptable" means a non-toxic material that does not interfere with the effectiveness of the biological activity of the active ingredient(s). The characteristics of the carrier will depend on the route of administration. The pharmaceutical composition of the invention may also contain cytokines, lymphokines, or other hematopoietic factors such as M-CSF, GM-CSF, TNF, IL-1, IL-2, IL-3, IL-4, IL-5, IL-6, IL-7, IL-8, IL-9, IL-10, IL-11, IL-12, IL-13, IL-14, IL-15, IFN, TNF0, TNF1, TNF2, G-CSF, Meg-CSF, thrombopoietin, stem cell factor, and erythropoietin. In further compositions, proteins of the invention may be combined with other agents beneficial to the treatment of the disease or disorder in question. These agents include various growth factors such as epidermal growth factor (EGF), platelet-derived growth factor (PDGF), transforming growth factors (TGF-α and TGF-β), insulin-like growth factor (IGF), as well as cytokines described herein.

The pharmaceutical composition may further contain other agents which either enhance the activity of the protein or other active ingredient or complement its activity or use in treatment. Such additional factors and/or agents may be included in the pharmaceutical composition to produce a synergistic effect with protein or other active ingredient of the invention, or to minimize side effects. Conversely, protein or other active ingredient of the present invention may be included in formulations of the particular clotting factor, cytokine, lymphokine, other hematopoietic factor, thrombolytic or anti-thrombotic factor, or anti-inflammatory agent to minimize side effects of the clotting factor, cytokine, lymphokine, other hematopoietic factor, thrombolytic or anti-thrombotic factor, or anti-inflammatory agent (such as IL-1Ra, IL-1 Hy1, IL-1 Hy2, anti-TNF, corticosteroids, immunosuppressive agents). A protein of the present invention may be active in multimers (e.g., heterodimers or homodimers) or complexes with itself or other proteins. As a result, pharmaceutical compositions of the invention may comprise a protein of the invention in such multimeric or complexed form.

As an alternative to being included in a pharmaceutical composition of the invention including a first protein, a second protein or a therapeutic agent may be concurrently administered with the first protein (e.g., at the same time, or at differing times provided that therapeutic concentrations of the combination of agents is achieved at the treatment site). Techniques for formulation and administration of the compounds of the instant application may be found in "Remington's Pharmaceutical Sciences," Mack Publishing Co., Easton, PA, latest edition. A therapeutically effective dose further refers to that amount of the compound sufficient to result in amelioration of symptoms, e.g., treatment, healing, prevention or amelioration of the relevant medical condition, or an increase in rate of treatment, healing, prevention or amelioration of such conditions. When applied to an individual active ingredient, administered

alone, a therapeutically effective dose refers to that ingredient alone. When applied to a combination, a therapeutically effective dose refers to combined amounts of the active ingredients that result in the therapeutic effect, whether administered in combination, serially or simultaneously.

In practicing the method of treatment or use of the present invention, a therapeutically effective amount of protein or other active ingredient of the present invention is administered to a mammal having a condition to be treated. Protein or other active ingredient of the present invention may be administered in accordance with the method of the invention either alone or in combination with other therapies such as treatments employing cytokines, lymphokines or other hematopoietic factors. When co- administered with one or more cytokines, lymphokines or other hematopoietic factors, protein or other active ingredient of the present invention may be administered either simultaneously with the cytokine(s), lymphokine(s), other hematopoietic factor(s), thrombolytic or anti-thrombotic factors, or sequentially. If administered sequentially, the attending physician will decide on the appropriate sequence of administering protein or other active ingredient of the present invention in combination with cytokine(s), lymphokine(s), other hematopoietic factor(s), thrombolytic or anti-thrombotic factors.

4.12.1 ROUTES OF ADMINISTRATION

Suitable routes of administration may, for example, include oral, rectal, transmucosal, or intestinal administration; parenteral delivery, including intramuscular, subcutaneous, intramedullary injections, as well as intrathecal, direct intraventricular, intravenous, intraperitoneal, intranasal, or intraocular injections. Administration of protein or other active ingredient of the present invention used in the pharmaceutical composition or to practice the method of the present invention can be carried out in a variety of conventional ways, such as oral ingestion, inhalation, topical application or cutaneous, subcutaneous, intraperitoneal, parenteral or intravenous injection. Intravenous administration to the patient is preferred.

Alternately, one may administer the compound in a local rather than systemic manner, for example, via injection of the compound directly into a arthritic joints or in fibrotic tissue; often in a depot or sustained release formulation. In order to prevent the scarring process frequently occurring as complication of glaucoma surgery, the compounds may be administered topically, for example, as eye drops. Furthermore, one may administer the drug in a targeted drug delivery system, for example, in a liposome coated with a specific antibody, targeting, for example, arthritic or fibrotic tissue. The liposomes will be targeted to and taken up selectively by the afflicted tissue.

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The polypeptides of the invention are administered by any route that delivers an effective dosage to the desired site of action. The determination of a suitable route of administration and an effective dosage for a particular indication is within the level of skill in the art. Preferably for wound treatment, one administers the therapeutic compound directly to the site. Suitable dosage ranges for the polypeptides of the invention can be extrapolated from these dosages or from similar studies in appropriate animal models. Dosages can then be adjusted as necessary by the clinician to provide maximal therapeutic benefit.

4.12.2 COMPOSITIONS/FORMULATIONS

Pharmaceutical compositions for use in accordance with the present invention thus may be formulated in a conventional manner using one or more physiologically acceptable carriers comprising excipients and auxiliaries which facilitate processing of the active compounds into preparations which can be used pharmaceutically. These pharmaceutical compositions may be manufactured in a manner that is itself known, e.g., by means of conventional mixing, dissolving, granulating, dragee-making, levigating, emulsifying, encapsulating, entrapping or lyophilizing processes. Proper formulation is dependent upon the route of administration chosen. When a therapeutically effective amount of protein or other active ingredient of the present invention is administered orally, protein or other active ingredient of the present invention will be in the form of a tablet, capsule, powder, solution or elixir. When administered in tablet form, the pharmaceutical composition of the invention may additionally contain a solid carrier such as a gelatin or an adjuvant. The tablet, capsule, and powder contain from about 5 to 95% protein or other active ingredient of the present invention, and preferably from about 25 to 90% protein or other active ingredient of the present invention. When administered in liquid form, a liquid carrier such as water, petroleum, oils of animal or plant origin such as peanut oil, mineral oil, soybean oil, or sesame oil, or synthetic oils may be added. The liquid form of the pharmaceutical composition may further contain physiological saline solution, dextrose or other saccharide solution, or glycols such as ethylene glycol, propylene glycol or polyethylene glycol. When administered in liquid form, the pharmaceutical composition contains from about 0.5 to 90% by weight of protein or other active ingredient of the present invention, and preferably from about 1 to 50% protein or other active ingredient of the present invention.

When a therapeutically effective amount of protein or other active ingredient of the present invention is administered by intravenous, cutaneous or subcutaneous injection, protein or other active ingredient of the present invention will be in the form of a pyrogen-free, parenterally acceptable aqueous solution. The preparation of such parenterally acceptable protein or other active ingredient solutions, having due regard to pH, isotonicity, stability, and the like, is within

the skill in the art. A preferred pharmaceutical composition for intravenous, cutaneous, or subcutaneous injection should contain, in addition to protein or other active ingredient of the present invention, an isotonic vehicle such as Sodium Chloride Injection, Ringer's Injection, Dextrose Injection, Dextrose and Sodium Chloride Injection, Lactated Ringer's Injection, or other vehicle as known in the art. The pharmaceutical composition of the present invention may also contain stabilizers, preservatives, buffers, antioxidants, or other additives known to those of skill in the art. For injection, the agents of the invention may be formulated in aqueous solutions, preferably in physiologically compatible buffers such as Hanks's solution, Ringer's solution, or physiological saline buffer. For transmucosal administration, penetrants appropriate to the barrier to be permeated are used in the formulation. Such penetrants are generally known in the art.

For oral administration, the compounds can be formulated readily by combining the active compounds with pharmaceutically acceptable carriers well known in the art. Such carriers enable the compounds of the invention to be formulated as tablets, pills, dragees, capsules, liquids, gels, syrups, slurries, suspensions and the like, for oral ingestion by a patient to be treated. Pharmaceutical preparations for oral use can be obtained from a solid excipient, optionally grinding a resulting mixture, and processing the mixture of granules, after adding suitable auxiliaries, if desired, to obtain tablets or dragee cores. Suitable excipients are, in particular, fillers such as sugars, including lactose, sucrose, mannitol, or sorbitol; cellulose preparations such as, for example, maize starch, wheat starch, rice starch, potato starch, gelatin, gum tragacanth, methyl cellulose, hydroxypropylmethyl-cellulose, sodium carboxymethylcellulose, and/or polyvinylpyrrolidone (PVP). If desired, disintegrating agents may be added, such as the cross-linked polyvinyl pyrrolidone, agar, or alginic acid or a salt thereof such as sodium alginate. Dragee cores are provided with suitable coatings. For this purpose, concentrated sugar solutions may be used, which may optionally contain gum arabic, talc, polyvinyl pyrrolidone, carbopol gel, polyethylene glycol, and/or titanium dioxide, lacquer solutions, and suitable organic solvents or solvent mixtures. Dyestuffs or pigments may be added to the tablets or dragee coatings for identification or to characterize different combinations of active compound doses.

Pharmaceutical preparations which can be used orally include push-fit capsules made of gelatin, as well as soft, sealed capsules made of gelatin and a plasticizer, such as glycerol or sorbitol. The push-fit capsules can contain the active ingredients in admixture with filler such as lactose, binders such as starches, and/or lubricants such as talc or magnesium stearate and, optionally, stabilizers. In soft capsules, the active compounds may be dissolved or suspended in suitable liquids, such as fatty oils, liquid paraffin, or liquid polyethylene glycols. In addition,

stabilizers may be added. All formulations for oral administration should be in dosages suitable for such administration. For buccal administration, the compositions may take the form of tablets or lozenges formulated in conventional manner.

For administration by inhalation, the compounds for use according to the present invention are conveniently delivered in the form of an aerosol spray presentation from pressurized packs or a nebuliser, with the use of a suitable propellant, e.g., dichlorodifluoromethane, trichlorofluoromethane, dichlorotetrafluoroethane, carbon dioxide or other suitable gas. In the case of a pressurized aerosol the dosage unit may be determined by providing a valve to deliver a metered amount. Capsules and cartridges of, e.g., gelatin for use in an inhaler or insufflator may be formulated containing a powder mix of the compound and a suitable powder base such as lactose or starch. The compounds may be formulated for parenteral administration by injection, e.g., by bolus injection or continuous infusion. Formulations for injection may be presented in unit dosage form, e.g., in ampules or in multi-dose containers, with an added preservative. The compositions may take such forms as suspensions, solutions or emulsions in oily or aqueous vehicles, and may contain formulatory agents such as suspending, stabilizing and/or dispersing agents.

Pharmaceutical formulations for parenteral administration include aqueous solutions of the active compounds in water-soluble form. Additionally, suspensions of the active compounds may be prepared as appropriate oily injection suspensions. Suitable lipophilic solvents or vehicles include fatty oils such as sesame oil, or synthetic fatty acid esters, such as ethyl oleate or triglycerides, or liposomes. Aqueous injection suspensions may contain substances which increase the viscosity of the suspension, such as sodium carboxymethyl cellulose, sorbitol, or dextran. Optionally, the suspension may also contain suitable stabilizers or agents which increase the solubility of the compounds to allow for the preparation of highly concentrated solutions. Alternatively, the active ingredient may be in powder form for constitution with a suitable vehicle, e.g., sterile pyrogen-free water, before use.

The compounds may also be formulated in rectal compositions such as suppositories or retention enemas, e.g., containing conventional suppository bases such as cocoa butter or other glycerides. In addition to the formulations described previously, the compounds may also be formulated as a depot preparation. Such long acting formulations may be administered by implantation (for example subcutaneously or intramuscularly) or by intramuscular injection. Thus, for example, the compounds may be formulated with suitable polymeric or hydrophobic materials (for example as an emulsion in an acceptable oil) or ion exchange resins, or as sparingly soluble derivatives, for example, as a sparingly soluble salt.

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A pharmaceutical carrier for the hydrophobic compounds of the invention is a co-solvent system comprising benzyl alcohol, a nonpolar surfactant, a water-miscible organic polymer, and an aqueous phase. The co-solvent system may be the VPD co-solvent system. VPD is a solution of 3% w/v benzyl alcohol, 8% w/v of the nonpolar surfactant polysorbate 80, and 65% w/v polyethylene glycol 300, made up to volume in absolute ethanol. The VPD co-solvent system (VPD:5W) consists of VPD diluted 1:1 with a 5% dextrose in water solution. This co-solvent system dissolves hydrophobic compounds well, and itself produces low toxicity upon systemic administration. Naturally, the proportions of a co-solvent system may be varied considerably without destroying its solubility and toxicity characteristics. Furthermore, the identity of the co-solvent components may be varied: for example, other low-toxicity nonpolar surfactants may be used instead of polysorbate 80; the fraction size of polyethylene glycol may be varied; other biocompatible polymers may replace polyethylene glycol, e.g. polyvinyl pyrrolidone; and other sugars or polysaccharides may substitute for dextrose. Alternatively, other delivery systems for hydrophobic pharmaceutical compounds may be employed. Liposomes and emulsions are well known examples of delivery vehicles or carriers for hydrophobic drugs. Certain organic solvents such as dimethylsulfoxide also may be employed, although usually at the cost of greater toxicity. Additionally, the compounds may be delivered using a sustained-release system, such as semipermeable matrices of solid hydrophobic polymers containing the therapeutic agent. Various types of sustained-release materials have been established and are well known by those skilled in the art. Sustained-release capsules may, depending on their chemical nature, release the compounds for a few weeks up to over 100 days. Depending on the chemical nature and the biological stability of the therapeutic reagent, additional strategies for protein or other active ingredient stabilization may be employed.

The pharmaceutical compositions also may comprise suitable solid or gel phase carriers or excipients. Examples of such carriers or excipients include but are not limited to calcium carbonate, calcium phosphate, various sugars, starches, cellulose derivatives, gelatin, and polymers such as polyethylene glycols. Many of the active ingredients of the invention may be provided as salts with pharmaceutically compatible counter ions. Such pharmaceutically acceptable base addition salts are those salts which retain the biological effectiveness and properties of the free acids and which are obtained by reaction with inorganic or organic bases such as sodium hydroxide, magnesium hydroxide, ammonia, trialkylamine, dialkylamine, monoalkylamine, dibasic amino acids, sodium acetate, potassium benzoate, triethanol amine and the like.

The pharmaceutical composition of the invention may be in the form of a complex of the protein(s) or other active ingredient(s) of present invention along with protein or peptide

antigens. The protein and/or peptide antigen will deliver a stimulatory signal to both B and T lymphocytes. B lymphocytes will respond to antigen through their surface immunoglobulin receptor. T lymphocytes will respond to antigen through the T cell receptor (TCR) following presentation of the antigen by MHC proteins. MHC and structurally related proteins including those encoded by class I and class II MHC genes on host cells will serve to present the peptide antigen(s) to T lymphocytes. The antigen components could also be supplied as purified MHC-peptide complexes alone or with co-stimulatory molecules that can directly signal T cells. Alternatively antibodies able to bind surface immunoglobulin and other molecules on B cells as well as antibodies able to bind the TCR and other molecules on T cells can be combined with the pharmaceutical composition of the invention.

The pharmaceutical composition of the invention may be in the form of a liposome in which protein of the present invention is combined, in addition to other pharmaceutically acceptable carriers, with amphipathic agents such as lipids which exist in aggregated form as micelles, insoluble monolayers, liquid crystals, or lamellar layers in aqueous solution. Suitable lipids for liposomal formulation include, without limitation, monoglycerides, diglycerides, sulfatides, lysolecithins, phospholipids, saponin, bile acids, and the like. Preparation of such liposomal formulations is within the level of skill in the art, as disclosed, for example, in U.S. Patent Nos. 4,235,871; 4,501,728; 4,837,028; and 4,737,323, all of which are incorporated herein by reference.

The amount of protein or other active ingredient of the present invention in the pharmaceutical composition of the present invention will depend upon the nature and severity of the condition being treated, and on the nature of prior treatments which the patient has undergone. Ultimately, the attending physician will decide the amount of protein or other active ingredient of the present invention with which to treat each individual patient. Initially, the attending physician will administer low doses of protein or other active ingredient of the present invention and observe the patient's response. Larger doses of protein or other active ingredient of the present invention may be administered until the optimal therapeutic effect is obtained for the patient, and at that point the dosage is not increased further. It is contemplated that the various pharmaceutical compositions used to practice the method of the present invention should contain about 0.01 µg to about 100 mg (preferably about 0.1 µg to about 10 mg, more preferably about 0.1 µg to about 1 mg) of protein or other active ingredient of the present invention per kg body weight. For compositions of the present invention which are useful for bone, cartilage, tendon or ligament regeneration, the therapeutic method includes administering the composition topically, systematically, or locally as an implant or device. When administered, the therapeutic composition for use in this invention is, of course, in a pyrogen-free, physiologically acceptable

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form. Further, the composition may desirably be encapsulated or injected in a viscous form for delivery to the site of bone, cartilage or tissue damage. Topical administration may be suitable for wound healing and tissue repair. Therapeutically useful agents other than a protein or other active ingredient of the invention which may also optionally be included in the composition as described above, may alternatively or additionally, be administered simultaneously or sequentially with the composition in the methods of the invention. Preferably for bone and/or cartilage formation, the composition would include a matrix capable of delivering the protein-containing or other active ingredient-containing composition to the site of bone and/or cartilage damage, providing a structure for the developing bone and cartilage and optimally capable of being resorbed into the body. Such matrices may be formed of materials presently in use for other implanted medical applications.

The choice of matrix material is based on biocompatibility, biodegradability, mechanical properties, cosmetic appearance and interface properties. The particular application of the compositions will define the appropriate formulation. Potential matrices for the compositions may be biodegradable and chemically defined calcium sulfate, tricalcium phosphate, hydroxyapatite, polylactic acid, polyglycolic acid and polyanhydrides. Other potential materials are biodegradable and biologically well-defined, such as bone or dermal collagen. Further matrices are comprised of pure proteins or extracellular matrix components. Other potential matrices are nonbiodegradable and chemically defined, such as sintered hydroxyapatite, bioglass, aluminates, or other ceramics. Matrices may be comprised of combinations of any of the above mentioned types of material, such as polylactic acid and hydroxyapatite or collagen and tricalcium phosphate. The bioceramics may be altered in composition, such as in calcium-aluminate-phosphate and processing to alter pore size, particle size, particle shape, and biodegradability. Presently preferred is a 50:50 (mole weight) copolymer of lactic acid and glycolic acid in the form of porous particles having diameters ranging from 150 to 800 microns. In some applications, it will be useful to utilize a sequestering agent, such as carboxymethyl cellulose or autologous blood clot, to prevent the protein compositions from disassociating from the matrix.

A preferred family of sequestering agents is cellulosic materials such as alkylcelluloses (including hydroxyalkylcelluloses), including methylcellulose, ethylcellulose, hydroxyethylcellulose, hydroxypropylcellulose, hydroxypropyl-methylcellulose, and carboxymethylcellulose, the most preferred being cationic salts of carboxymethylcellulose (CMC). Other pref rred sequestering agents include hyaluronic acid, sodium alginate, poly(ethylene glycol), polyoxyethylene oxide, carboxyvinyl polymer and poly(vinyl alcohol). The amount of sequestering agent useful herein is 0.5-20 wt %, preferably 1-10 wt % based on 72

total formulation weight, which represents the amount necessary to prevent desorption of the protein from the polymer matrix and to provide appropriate handling of the composition, yet not so much that the progenitor cells are prevented from infiltrating the matrix, thereby providing the protein the opportunity to assist the osteogenic activity of the progenitor cells. In further compositions, proteins or other active ingredients of the invention may be combined with other agents beneficial to the treatment of the bone and/or cartilage defect, wound, or tissue in question. These agents include various growth factors such as epidermal growth factor (EGF), platelet derived growth factor (PDGF), transforming growth factors (TGF-α and TGF-β), and insulin-like growth factor (IGF).

The therapeutic compositions are also presently valuable for veterinary applications. Particularly domestic animals and thoroughbred horses, in addition to humans, are desired patients for such treatment with proteins or other active ingredients of the present invention. The dosage regimen of a protein-containing pharmaceutical composition to be used in tissue regeneration will be determined by the attending physician considering various factors which modify the action of the proteins, e.g., amount of tissue weight desired to be formed, the site of damage, the condition of the damaged tissue, the size of a wound, type of damaged tissue (e.g., bone), the patient's age, sex, and diet, the severity of any infection, time of administration and other clinical factors. The dosage may vary with the type of matrix used in the reconstitution and with inclusion of other proteins in the pharmaceutical composition. For example, the addition of other known growth factors, such as IGF I (insulin like growth factor I), to the final composition, may also effect the dosage. Progress can be monitored by periodic assessment of tissue/bone growth and/or repair, for example, X-rays, histomorphometric determinations and tetracycline labeling.

Polynucleotides of the present invention can also be used for gene therapy. Such polynucleotides can be introduced either in vivo or ex vivo into cells for expression in a mammalian subject. Polynucleotides of the invention may also be administered by other known methods for introduction of nucleic acid into a cell or organism (including, without limitation, in the form of viral vectors or naked DNA). Cells may also be cultured ex vivo in the presence of proteins of the present invention in order to proliferate or to produce a desired effect on or activity in such cells. Treated cells can then be introduced in vivo for therapeutic purposes.

4.12.3 EFFECTIVE DOSAGE

Pharmaceutical compositions suitable for use in the present invention include compositions wherein the active ingredients are contained in an effective amount to achieve its intended purpose. More specifically, a therapeutically effective amount means an amount

effective to prevent development of or to alleviate the existing symptoms of the subject being treated. Determination of the effective amount is well within the capability of those skilled in the art, especially in light of the detailed disclosure provided herein. For any compound used in the method of the invention, the therapeutically effective dose can be estimated initially from appropriate in vitro assays. For example, a dose can be formulated in animal models to achieve a circulating concentration range that can be used to more accurately determine useful doses in humans. For example, a dose can be formulated in animal models to achieve a circulating concentration range that includes the IC₅₀ as determined in cell culture (*i.e.*, the concentration of the test compound which achieves a half-maximal inhibition of the protein's biological activity). Such information can be used to more accurately determine useful doses in humans.

A therapeutically effective dose refers to that amount of the compound that results in amelioration of symptoms or a prolongation of survival in a patient. Toxicity and therapeutic efficacy of such compounds can be determined by standard pharmaceutical procedures in cell cultures or experimental animals, e.g., for determining the LD50 (the dose lethal to 50% of the population) and the ED₅₀ (the dose therapeutically effective in 50% of the population). The dose ratio between toxic and therapeutic effects is the therapeutic index and it can be expressed as the ratio between LD₅₀ and ED₅₀. Compounds which exhibit high therapeutic indices are preferred. The data obtained from these cell culture assays and animal studies can be used in formulating a range of dosage for use in human. The dosage of such compounds lies preferably within a range of circulating concentrations that include the ED50 with little or no toxicity. The dosage may vary within this range depending upon the dosage form employed and the route of administration utilized. The exact formulation, route of administration and dosage can be chosen by the individual physician in view of the patient's condition. See, e.g., Fingl et al., 1975, in "The Pharmacological Basis of Therapeutics", Ch. 1 p.1. Dosage amount and interval may be adjusted individually to provide plasma levels of the active moiety which are sufficient to maintain the desired effects, or minimal effective concentration (MEC). The MEC will vary for each compound but can be estimated from in vitro data. Dosages necessary to achieve the MEC will depend on individual characteristics and route of administration. However, HPLC assays or bioassays can be used to determine plasma concentrations.

Dosage intervals can also be determined using MEC value. Compounds should be administered using a regimen which maintains plasma levels above the MEC for 10-90% of the time, preferably between 30-90% and most preferably between 50-90%. In cases of local administration or selective uptake, the effective local concentration of the drug may not be related to plasma concentration.

An exemplary dosage regimen for polypeptides or other compositions of the invention will be in the range of about 0.01 μ g/kg to 100 mg/kg of body weight daily, with the preferred dose being about 0.1 μ g/kg to 25 mg/kg of patient body weight daily, varying in adults and children. Dosing may be once daily, or equivalent doses may be delivered at longer or shorter intervals.

The amount of composition administered will, of course, be dependent on the subject being treated, on the subject's age and weight, the severity of the affliction, the manner of administration and the judgment of the prescribing physician.

4.12.4 PACKAGING

The compositions may, if desired, be presented in a pack or dispenser device which may contain one or more unit dosage forms containing the active ingredient. The pack may, for example, comprise metal or plastic foil, such as a blister pack. The pack or dispenser device may be accompanied by instructions for administration. Compositions comprising a compound of the invention formulated in a compatible pharmaceutical carrier may also be prepared, placed in an appropriate container, and labeled for treatment of an indicated condition.

4.13 ANTIBODIES

Also included in the invention are antibodies to proteins, or fragments of proteins of the invention. The term "antibody" as used herein refers to immunoglobulin molecules and immunologically active portions of immunoglobulin (Ig) molecules, i.e., molecules that contain an antigen binding site that specifically binds (immunoreacts with) an antigen. Such antibodies include, but are not limited to, polyclonal, monoclonal, chimeric, single chain, F_{ab} , F_{ab} , and $F_{(ab)}$ fragments, and an F_{ab} expression library. In general, an antibody molecule obtained from humans relates to any of the classes IgG, IgM, IgA, IgE and IgD, which differ from one another by the nature of the heavy chain present in the molecule. Certain classes have subclasses as well, such as IgG₁, IgG₂, and others. Furthermore, in humans, the light chain may be a kappa chain or a lambda chain. Reference herein to antibodies includes a reference to all such classes, subclasses and types of human antibody species.

An isolated related protein of the invention may be intended to serve as an antigen, or a portion or fragment thereof, and additionally can be used as an immunogen to generate antibodies that immunospecifically bind the antigen, using standard techniques for polyclonal and monoclonal antibody preparation. The full-length protein can be used or, alternatively, the invention provides antigenic peptide fragments of the antigen for use as immunogens. An antigenic peptide fragment comprises at least 6 amino acid residues of the amino acid sequence

of the full length protein, such as the amino acid sequences shown in SEQ ID NO: 445-888, and encompasses an epitope thereof such that an antibody raised against the peptide forms a specific immune complex with the full length protein or with any fragment that contains the epitope. Preferably, the antigenic peptide comprises at least 10 amino acid residues, or at least 15 amino acid residues, or at least 20 amino acid residues, or at least 30 amino acid residues. Preferred epitopes encompassed by the antigenic peptide are regions of the protein that are located on its surface; commonly these are hydrophilic regions.

In certain embodiments of the invention, at least one epitope encompassed by the antigenic peptide is a region of -related protein that is located on the surface of the protein, e.g., a hydrophilic region. A hydrophobicity analysis of the human related protein sequence will indicate which regions of a related protein are particularly hydrophilic and, therefore, are likely to encode surface residues useful for targeting antibody production. As a means for targeting antibody production, hydropathy plots showing regions of hydrophilicity and hydrophobicity may be generated by any method well known in the art, including, for example, the Kyte Doolittle or the Hopp Woods methods, either with or without Fourier transformation. See, e.g., Hopp and Woods, 1981, Proc. Nat. Acad. Sci. USA 78: 3824-3828; Kyte and Doolittle 1982, J. Mol. Biol. 157: 105-142, each of which is incorporated herein by reference in its entirety. Antibodies that are specific for one or more domains within an antigenic protein, or derivatives, fragments, analogs or homologs thereof, are also provided herein.

A protein of the invention, or a derivative, fragment, analog, homolog or ortholog thereof, may be utilized as an immunogen in the generation of antibodies that immunospecifically bind these protein components.

Various procedures known within the art may be used for the production of polyclonal or monoclonal antibodies directed against a protein of the invention, or against derivatives, fragments, analogs homologs or orthologs thereof (see, for example, Antibodies: A Laboratory Manual, Harlow E, and Lane D, 1988, Cold Spring Harbor Laboratory Press, Cold Spring Harbor, NY, incorporated herein by reference). Some of these antibodies are discussed below.

4.13.1 POLYCLONAL ANTIBODIES

For the production of polyclonal antibodies, various suitable host animals (e.g., rabbit, goat, mouse or other mammal) may be immunized by one or more injections with the native protein, a synthetic variant thereof, or a derivative of the foregoing. An appropriate immunogenic preparation can contain, for example, the naturally occurring immunogenic protein, a chemically synthesized polypeptide representing the immunogenic protein, or a recombinantly expressed immunogenic protein. Furthermore, the protein may be conjugated to

a second protein known to be immunogenic in the mammal being immunized. Examples of such immunogenic proteins include but are not limited to keyhole limpet hemocyanin, serum albumin, bovine thyroglobulin, and soybean trypsin inhibitor. The preparation can further include an adjuvant. Various adjuvants used to increase the immunological response include, but are not limited to, Freund's (complete and incomplete), mineral gels (e.g., aluminum hydroxide), surface active substances (e.g., lysolecithin, pluronic polyols, polyanions, peptides, oil emulsions, dinitrophenol, etc.), adjuvants usable in humans such as Bacille Calmette-Guerin and Corynebacterium parvum, or similar immunostimulatory agents. Additional examples of adjuvants which can be employed include MPL-TDM adjuvant (monophosphoryl Lipid A, synthetic trehalose dicorynomycolate).

The polyclonal antibody molecules directed against the immunogenic protein can be isolated from the mammal (e.g., from the blood) and further purified by well known techniques, such as affinity chromatography using protein A or protein G, which provide primarily the IgG fraction of immune serum. Subsequently, or alternatively, the specific antigen which is the target of the immunoglobulin sought, or an epitope thereof, may be immobilized on a column to purify the immune specific antibody by immunoaffinity chromatography. Purification of immunoglobulins is discussed, for example, by D. Wilkinson (The Scientist, published by The Scientist, Inc., Philadelphia PA, Vol. 14, No. 8 (April 17, 2000), pp. 25-28).

4.13.2 MONOCLONAL ANTIBODIES

The term "monoclonal antibody" (MAb) or "monoclonal antibody composition", as used herein, refers to a population of antibody molecules that contain only one molecular species of antibody molecule consisting of a unique light chain gene product and a unique heavy chain gene product. In particular, the complementarity determining regions (CDRs) of the monoclonal antibody are identical in all the molecules of the population. MAbs thus contain an antigen binding site capable of immunoreacting with a particular epitope of the antigen characterized by a unique binding affinity for it.

Monoclonal antibodies can be prepared using hybridoma methods, such as those described by Kohler and Milstein, Nature, 256:495 (1975). In a hybridoma method, a mouse, hamster, or other appropriate host animal, is typically immunized with an immunizing agent to elicit lymphocytes that produce or are capable of producing antibodies that will specifically bind to the immunizing agent. Alternatively, the lymphocytes can be immunized in vitro. The immunizing agent will typically include the protein antigen, a fragment thereof or a fusion protein thereof. Generally, either peripheral blood lymphocytes are used if cells of human origin are desired, or spleen cells or lymph node cells are used if non-human mammalian sources are

desired. The lymphocytes are then fused with an immortalized cell line using a suitable fusing agent, such as polyethylene glycol, to form a hybridoma cell (Goding, Monoclonal Antibodies: Principles and Practice, Academic Press, (1986) pp. 59-103). Immortalized cell lines are usually transformed mammalian cells, particularly myeloma cells of rodent, bovine and human origin. Usually, rat or mouse myeloma cell lines are employed. The hybridoma cells can be cultured in a suitable culture medium that preferably contains one or more substances that inhibit the growth or survival of the unfused, immortalized cells. For example, if the parental cells lack the enzyme hypoxanthine guanine phosphoribosyl transferase (HGPRT or HPRT), the culture medium for the hybridomas typically will include hypoxanthine, aminopterin, and thymidine ("HAT medium"), which substances prevent the growth of HGPRT-deficient cells.

Preferred immortalized cell lines are those that fuse efficiently, support stable high level expression of antibody by the selected antibody-producing cells, and are sensitive to a medium such as HAT medium. More preferred immortalized cell lines are murine myeloma lines, which can be obtained, for instance, from the Salk Institute Cell Distribution Center, San Diego, California and the American Type Culture Collection, Manassas, Virginia. Human myeloma and mouse-human heteromyeloma cell lines also have been described for the production of human monoclonal antibodies (Kozbor, J. Immunol., 133:3001 (1984); Brodeur et al., Monoclonal Antibody Production Techniques and Applications, Marcel Dekker, Inc., New York, (1987) pp. 51-63).

The culture medium in which the hybridoma cells are cultured can then be assayed for the presence of monoclonal antibodies directed against the antigen. Preferably, the binding specificity of monoclonal antibodies produced by the hybridoma cells is determined by immunoprecipitation or by an in vitro binding assay, such as radioimmunoassay (RIA) or enzyme-linked immunoabsorbent assay (ELISA). Such techniques and assays are known in the art. The binding affinity of the monoclonal antibody can, for example, be determined by the Scatchard analysis of Munson and Pollard, <u>Anal. Biochem.</u>, 107:220 (1980). Preferably, antibodies having a high degree of specificity and a high binding affinity for the target antigen are isolated.

After the desired hybridoma cells are identified, the clones can be subcloned by limiting dilution procedures and grown by standard methods. Suitable culture media for this purpose include, for example, Dulbecco's Modified Eagle's Medium and RPMI-1640 medium.

Alternatively, the hybridoma cells can be grown in vivo as ascites in a mammal.

The monoclonal antibodies secreted by the subclones can be isolated or purified from the culture medium or ascites fluid by conventional immunoglobulin purification procedures such as, for

example, protein A-Sepharose, hydroxylapatite chromatography, gel electrophoresis, dialysis, or affinity chromatography.

The monoclonal antibodies can also be made by recombinant DNA methods, such as those described in U.S. Patent No. 4,816,567. DNA encoding the monoclonal antibodies of the invention can be readily isolated and sequenced using conventional procedures (e.g., by using oligonucleotide probes that are capable of binding specifically to genes encoding the heavy and light chains of murine antibodies). The hybridoma cells of the invention serve as a preferred source of such DNA. Once isolated, the DNA can be placed into expression vectors, which are then transfected into host cells such as simian COS cells, Chinese hamster ovary (CHO) cells, or myeloma cells that do not otherwise produce immunoglobulin protein, to obtain the synthesis of monoclonal antibodies in the recombinant host cells. The DNA also can be modified, for example, by substituting the coding sequence for human heavy and light chain constant domains in place of the homologous murine sequences (U.S. Patent No. 4,816,567; Morrison, Nature 368, 812-13 (1994)) or by covalently joining to the immunoglobulin coding sequence all or part of the coding sequence for a non-immunoglobulin polypeptide. Such a non-immunoglobulin polypeptide can be substituted for the constant domains of an antibody of the invention, or can be substituted for the variable domains of one antigen-combining site of an antibody of the invention to create a chimeric bivalent antibody.

4.13.3 HUMANIZED ANTIBODIES

The antibodies directed against the protein antigens of the invention can further comprise humanized antibodies or human antibodies. These antibodies are suitable for administration to humans without engendering an immune response by the human against the administered immunoglobulin. Humanized forms of antibodies are chimeric immunoglobulins, immunoglobulin chains or fragments thereof (such as Fv, Fab, Fab', F(ab')2 or other antigenbinding subsequences of antibodies) that are principally comprised of the sequence of a human immunoglobulin, and contain minimal sequence derived from a non-human immunoglobulin. Humanization can be performed following the method of Winter and co-workers (Jones et al., Nature, 321:522-525 (1986); Riechmann et al., Nature, 332:323-327 (1988); Verhoeyen et al., Science, 239:1534-1536 (1988)), by substituting rodent CDRs or CDR sequences for the corresponding sequences of a human antibody. (See also U.S. Patent No. 5,225,539.) In some instances, Fv framework residues of the human immunoglobulin are replaced by corresponding non-human residues. Humanized antibodies can also comprise residues which are found neither in the recipient antibody nor in the imported CDR or framework sequences. In general, the humanized antibody will comprise substantially all of at least one, and typically two, variable

domains, in which all or substantially all of the CDR regions correspond to those of a non-human immunoglobulin and all or substantially all of the framework regions are those of a human immunoglobulin consensus sequence. The humanized antibody optimally also will comprise at least a portion of an immunoglobulin constant region (Fc), typically that of a human immunoglobulin (Jones et al., 1986; Riechmann et al., 1988; and Presta, <u>Curr. Op. Struct. Biol.</u>, 2:593-596 (1992)).

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4.13.4 HUMAN ANTIBODIES

Fully human antibodies relate to antibody molecules in which essentially the entire sequences of both the light chain and the heavy chain, including the CDRs, arise from human genes. Such antibodies are termed "human antibodies", or "fully human antibodies" herein. Human monoclonal antibodies can be prepared by the trioma technique; the human B-cell hybridoma technique (see Kozbor, et al., 1983 Immunol Today 4: 72) and the EBV hybridoma technique to produce human monoclonal antibodies (see Cole, et al., 1985 In: MONOCLONAL ANTIBODIES AND CANCER THERAPY, Alan R. Liss, Inc., pp. 77-96). Human monoclonal antibodies may be utilized in the practice of the present invention and may be produced by using human hybridomas (see Cote, et al., 1983. Proc Natl Acad Sci USA 80: 2026-2030) or by transforming human B-cells with Epstein Barr Virus in vitro (see Cole, et al., 1985 In: MONOCLONAL ANTIBODIES AND CANCER THERAPY, Alan R. Liss, Inc., pp. 77-96).

In addition, human antibodies can also be produced using additional techniques, including phage display libraries (Hoogenboom and Winter, J. Mol. Biol., 227:381 (1991); Marks et al., J. Mol. Biol., 222:581 (1991)). Similarly, human antibodies can be made by introducing human immunoglobulin loci into transgenic animals, e.g., mice in which the endogenous immunoglobulin genes have been partially or completely inactivated. Upon challenge, human antibody production is observed, which closely resembles that seen in humans in all respects, including gene rearrangement, assembly, and antibody repertoire. This approach is described, for example, in U.S. Patent Nos. 5,545,807; 5,545,806; 5,569,825; 5,625,126; 5,633,425; 5,661,016, and in Marks et al. (Bio/Technology 10, 779-783 (1992)); Lonberg et al. (Nature 368 856-859 (1994)); Morrison (Nature 368, 812-13 (1994)); Fishwild et al., (Nature Biotechnology 14, 845-51 (1996)); Neuberger (Nature Biotechnology 14, 826 (1996)); and Lonberg and Huszar (Intern. Rev. Immunol. 13 65-93 (1995)).

Human antibodies may additionally be produced using transgenic nonhuman animals which are modified so as to produce fully human antibodies rather than the animal's endogenous antibodies in response to challenge by an antigen. (See PCT publication WO94/02602). The endogenous genes encoding the heavy and light immunoglobulin chains in the nonhuman host

have been incapacitated, and active loci encoding human heavy and light chain immunoglobulins are inserted into the host's genome. The human genes are incorporated, for example, using yeast artificial chromosomes containing the requisite human DNA segments. An animal which provides all the desired modifications is then obtained as progeny by crossbreeding intermediate transgenic animals containing fewer than the full complement of the modifications. The preferred embodiment of such a nonhuman animal is a mouse, and is termed the Xenomouse TM as disclosed in PCT publications WO 96/33735 and WO 96/34096. This animal produces B cells which secrete fully human immunoglobulins. The antibodies can be obtained directly from the animal after immunization with an immunogen of interest, as, for example, a preparation of a polyclonal antibody, or alternatively from immortalized B cells derived from the animal, such as hybridomas producing monoclonal antibodies. Additionally, the genes encoding the immunoglobulins with human variable regions can be recovered and expressed to obtain the antibodies directly, or can be further modified to obtain analogs of antibodies such as, for example, single chain Fv molecules.

An example of a method of producing a nonhuman host, exemplified as a mouse, lacking expression of an endogenous immunoglobulin heavy chain is disclosed in U.S. Patent No. 5,939,598. It can be obtained by a method including deleting the J segment genes from at least one endogenous heavy chain locus in an embryonic stem cell to prevent rearrangement of the locus and to prevent formation of a transcript of a rearranged immunoglobulin heavy chain locus, the deletion being effected by a targeting vector containing a gene encoding a selectable marker; and producing from the embryonic stem cell a transgenic mouse whose somatic and germ cells contain the gene encoding the selectable marker.

A method for producing an antibody of interest, such as a human antibody, is disclosed in U.S. Patent No. 5,916,771. It includes introducing an expression vector that contains a nucleotide sequence encoding a heavy chain into one mammalian host cell in culture, introducing an expression vector containing a nucleotide sequence encoding a light chain into another mammalian host cell, and fusing the two cells to form a hybrid cell. The hybrid cell expresses an antibody containing the heavy chain and the light chain.

In a further improvement on this procedure, a method for identifying a clinically relevant epitope on an immunogen, and a correlative method for selecting an antibody that binds immunospecifically to the relevant epitope with high affinity, are disclosed in PCT publication WO 99/53049.

4.13.5 Fab FRAGMENTS AND SINGLE CHAIN ANTIBODIES

According to the invention, techniques can be adapted for the production of single-chain antibodies specific to an antigenic protein of the invention (see e.g., U.S. Patent No. 4,946,778). In addition, methods can be adapted for the construction of F_{ab} expression libraries (see e.g., Huse, et al., 1989 Science 246: 1275-1281) to allow rapid and effective identification of monoclonal F_{ab} fragments with the desired specificity for a protein or derivatives, fragments, analogs or homologs thereof. Antibody fragments that contain the idiotypes to a protein antigen may be produced by techniques known in the art including, but not limited to: (i) an $F_{(ab')2}$ fragment produced by pepsin digestion of an antibody molecule; (ii) an F_{ab} fragment generated by reducing the disulfide bridges of an $F_{(ab')2}$ fragment; (iii) an F_{ab} fragment generated by the treatment of the antibody molecule with papain and a reducing agent and (iv) F_{v} fragments.

4.13.6 BISPECIFIC ANTIBODIES

Bispecific antibodies are monoclonal, preferably human or humanized, antibodies that have binding specificities for at least two different antigens. In the present case, one of the binding specificities is for an antigenic protein of the invention. The second binding target is any other antigen, and advantageously is a cell-surface protein or receptor or receptor subunit.

Methods for making bispecific antibodies are known in the art. Traditionally, the recombinant production of bispecific antibodies is based on the co-expression of two immunoglobulin heavy-chain/light-chain pairs, where the two heavy chains have different specificities (Milstein and Cuello, Nature, 305:537-539 (1983)). Because of the random assortment of immunoglobulin heavy and light chains, these hybridomas (quadromas) produce a potential mixture of ten different antibody molecules, of which only one has the correct bispecific structure. The purification of the correct molecule is usually accomplished by affinity chromatography steps. Similar procedures are disclosed in WO 93/08829, published 13 May 1993, and in Traumecker et al., 1991 EMBO J., 10:3655-3659.

Antibody variable domains with the desired binding specificities (antibody-antigen combining sites) can be fused to immunoglobulin constant domain sequences. The fusion preferably is with an immunoglobulin heavy-chain constant domain, comprising at least part of the hinge, CH2, and CH3 regions. It is preferred to have the first heavy-chain constant region (CH1) containing the site necessary for light-chain binding present in at least one of the fusions. DNAs encoding the immunoglobulin heavy-chain fusions and, if desired, the immunoglobulin light chain, are inserted into separate expression vectors, and are co-transfected into a suitable host organism. For further details of generating bispecific antibodies see, for example, Suresh et al., Methods in Enzymology, 121:210 (1986).

According to another approach described in WO 96/27011, the interface between a pair of antibody molecules can be engineered to maximize the percentage of heterodimers which are recovered from recombinant cell culture. The preferred interface comprises at least a part of the CH3 region of an antibody constant domain. In this method, one or more small amino acid side chains from the interface of the first antibody molecule are replaced with larger side chains (e.g. tyrosine or tryptophan). Compensatory "cavities" of identical or similar size to the large side chain(s) are created on the interface of the second antibody molecule by replacing large amino acid side chains with smaller ones (e.g. alanine or threonine). This provides a mechanism for increasing the yield of the heterodimer over other unwanted end-products such as homodimers.

Bispecific antibodies can be prepared as full length antibodies or antibody fragments (e.g. F(ab')₂ bispecific antibodies). Techniques for generating bispecific antibodies from antibody fragments have been described in the literature. For example, bispecific antibodies can be prepared using chemical linkage. Brennan et al., Science 229:81 (1985) describe a procedure wherein intact antibodies are proteolytically cleaved to generate F(ab')₂ fragments. These fragments are reduced in the presence of the dithiol complexing agent sodium arsenite to stabilize vicinal dithiols and prevent intermolecular disulfide formation. The Fab' fragments generated are then converted to thionitrobenzoate (TNB) derivatives. One of the Fab'-TNB derivatives is then reconverted to the Fab'-thiol by reduction with mercaptoethylamine and is mixed with an equimolar amount of the other Fab'-TNB derivative to form the bispecific antibody. The bispecific antibodies produced can be used as agents for the selective immobilization of enzymes.

Additionally, Fab' fragments can be directly recovered from E. coli and chemically coupled to form bispecific antibodies. Shalaby et al., J. Exp. Med. 175:217-225 (1992) describe the production of a fully humanized bispecific antibody F(ab')₂ molecule. Each Fab' fragment was separately secreted from E. coli and subjected to directed chemical coupling in vitro to form the bispecific antibody. The bispecific antibody thus formed was able to bind to cells overexpressing the ErbB2 receptor and normal human T cells, as well as trigger the lytic activity of human cytotoxic lymphocytes against human breast tumor targets.

Various techniques for making and isolating bispecific antibody fragments directly from recombinant cell culture have also been described. For example, bispecific antibodies have been produced using leucine zippers. Kostelny et al., J. Immunol. 148(5):1547-1553 (1992). The leucine zipper peptides from the Fos and Jun proteins were linked to the Fab' portions of two different antibodies by gene fusion. The antibody homodimers were reduced at the hinge region to form monomers and then re-oxidized to form the antibody heterodimers. This method can also be utilized for the production of antibody homodimers. The "diabody" technology

described by Hollinger et al., <u>Proc. Natl. Acad. Sci. USA</u> 90:6444-6448 (1993) has provided an alternative mechanism for making bispecific antibody fragments. The fragments comprise a heavy-chain variable domain (V_L) by a linker which is too short to allow pairing between the two domains on the same chain. Accordingly, the V_H and V_L domains of one fragment are forced to pair with the complementary V_L and V_H domains of another fragment, thereby forming two antigen-binding sites. Another strategy for making bispecific antibody fragments by the use of single-chain Fv (sFv) dimers has also been reported. See, Gruber et al., <u>J. Immunol.</u> 152:5368 (1994).

Antibodies with more than two valencies are contemplated. For example, trispecific antibodies can be prepared. Tutt et al., J. Immunol. 147:60 (1991).

Exemplary bispecific antibodies can bind to two different epitopes, at least one of which originates in the protein antigen of the invention. Alternatively, an anti-antigenic arm of an immunoglobulin molecule can be combined with an arm which binds to a triggering molecule on a leukocyte such as a T-cell receptor molecule (e.g. CD2, CD3, CD28, or B7), or Fc receptors for IgG (FcyR), such as FcyRI (CD64), FcyRII (CD32) and FcyRIII (CD16) so as to focus cellular defense mechanisms to the cell expressing the particular antigen. Bispecific antibodies can also be used to direct cytotoxic agents to cells which express a particular antigen. These antibodies possess an antigen-binding arm and an arm which binds a cytotoxic agent or a radionuclide chelator, such as EOTUBE, DPTA, DOTA, or TETA. Another bispecific antibody of interest binds the protein antigen described herein and further binds tissue factor (TF).

4.13.7 HETEROCONJUGATE ANTIBODIES

Heteroconjugate antibodies are also within the scope of the present invention. Heteroconjugate antibodies are composed of two covalently joined antibodies. Such antibodies have, for example, been proposed to target immune system cells to unwanted cells (U.S. Patent No. 4,676,980), and for treatment of HIV infection (WO 91/00360; WO 92/200373; EP 03089). It is contemplated that the antibodies can be prepared in vitro using known methods in synthetic protein chemistry, including those involving crosslinking agents. For example, immunotoxins can be constructed using a disulfide exchange reaction or by forming a thioether bond. Examples of suitable reagents for this purpose include iminothiolate and methyl-4-mercaptobutyrimidate and those disclosed, for example, in U.S. Patent No. 4,676,980.

4.13.8 EFFECTOR FUNCTION ENGINEERING

It can be desirable to modify the antibody of the invention with respect to effector function, so as to enhance, e.g., the effectiveness of the antibody in treating cancer. For example, cysteine

residue(s) can be introduced into the Fc region, thereby allowing interchain disulfide bond formation in this region. The homodimeric antibody thus generated can have improved internalization capability and/or increased complement-mediated cell killing and antibody-dependent cellular cytotoxicity (ADCC). See Caron et al., J. Exp Med., 176: 1191-1195 (1992) and Shopes, J. Immunol., 148: 2918-2922 (1992). Homodimeric antibodies with enhanced antitumor activity can also be prepared using heterobifunctional cross-linkers as described in Wolff et al. Cancer Research, 53: 2560-2565 (1993). Alternatively, an antibody can be engineered that has dual Fc regions and can thereby have enhanced complement lysis and ADCC capabilities. See Stevenson et al., Anti-Cancer Drug Design, 3: 219-230 (1989).

4.13.9 IMMUNOCONJUGATES

The invention also pertains to immunoconjugates comprising an antibody conjugated to a cytotoxic agent such as a chemotherapeutic agent, toxin (e.g., an enzymatically active toxin of bacterial, fungal, plant, or animal origin, or fragments thereof), or a radioactive isotope (i.e., a radioconjugate).

Chemotherapeutic agents useful in the generation of such immunoconjugates have been described above. Enzymatically active toxins and fragments thereof that can be used include diphtheria A chain, nonbinding active fragments of diphtheria toxin, exotoxin A chain (from Pseudomonas aeruginosa), ricin A chain, abrin A chain, modeccin A chain, alpha-sarcin, Aleurites fordii proteins, dianthin proteins, Phytolaca americana proteins (PAPI, PAPII, and PAP-S), momordica charantia inhibitor, curcin, crotin, sapaonaria officinalis inhibitor, gelonin, mitogellin, restrictocin, phenomycin, enomycin, and the tricothecenes. A variety of radionuclides are available for the production of radioconjugated antibodies. Examples include ²¹²Bi, ¹³¹I, ¹³¹In, ⁹⁰Y, and ¹⁸⁶Re.

Conjugates of the antibody and cytotoxic agent are made using a variety of bifunctional protein-coupling agents such as N-succinimidyl-3-(2-pyridyldithiol) propionate (SPDP), iminothiolane (IT), bifunctional derivatives of imidoesters (such as dimethyl adipimidate HCL), active esters (such as disuccinimidyl suberate), aldehydes (such as glutareldehyde), bis-azido compounds (such as bis (p-azidobenzoyl) hexanediamine), bis-diazonium derivatives (such as bis-(p-diazoniumbenzoyl)-ethylenediamine), diisocyanates (such as tolyene 2,6-diisocyanate), and bis-active fluorine compounds (such as 1,5-difluoro-2,4-dinitrobenzene). For example, a ricin immunotoxin can be prepared as described in Vitetta et al., Science, 238: 1098 (1987). Carbon-14-labeled 1-isothiocyanatobenzyl-3-methyldiethylene triaminepentaacetic acid (MX-DTPA) is an exemplary chelating agent for conjugation of radionucleotide to the antibody. See WO94/11026.

In another embodiment, the antibody can be conjugated to a "receptor" (such as streptavidin) for utilization in tumor pretargeting wherein the antibody-receptor conjugate is administered to the patient, followed by removal of unbound conjugate from the circulation using a clearing agent and then administration of a "ligand" (e.g., avidin) that is in turn conjugated to a cytotoxic agent.

4.14 COMPUTER READABLE SEQUENCES

In one application of this embodiment, a nucleotide sequence of the present invention can be recorded on computer readable media. As used herein, "computer readable media" refers to any medium which can be read and accessed directly by a computer. Such media include, but are not limited to: magnetic storage media, such as floppy discs, hard disc storage medium, and magnetic tape; optical storage media such as CD-ROM; electrical storage media such as RAM and ROM; and hybrids of these categories such as magnetic/optical storage media. A skilled artisan can readily appreciate how any of the presently known computer readable mediums can be used to create a manufacture comprising computer readable medium having recorded thereon a nucleotide sequence of the present invention. As used herein, "recorded" refers to a process for storing information on computer readable medium. A skilled artisan can readily adopt any of the presently known methods for recording information on computer readable medium to generate manufactures comprising the nucleotide sequence information of the present invention.

A variety of data storage structures are available to a skilled artisan for creating a computer readable medium having recorded thereon a nucleotide sequence of the present invention. The choice of the data storage structure will generally be based on the means chosen to access the stored information. In addition, a variety of data processor programs and formats can be used to store the nucleotide sequence information of the present invention on computer readable medium. The sequence information can be represented in a word processing text file, formatted in commercially-available software such as WordPerfect and Microsoft Word, or represented in the form of an ASCII file, stored in a database application, such as DB2, Sybase, Oracle, or the like. A skilled artisan can readily adapt any number of data processor structuring formats (e.g. text file or database) in order to obtain computer readable medium having recorded thereon the nucleotide sequence information of the present invention.

By providing any of the nucleotide sequences SEQ ID NO: 1-444 or a representative fragment thereof; or a nucleotide sequence at least 95% identical to any of the nucleotide sequences of SEQ ID NO: 1-444 in computer readable form, a skilled artisan can routinely access the sequence information for a variety of purposes. Computer software is publicly available which allows a skilled artisan to access sequence information provided in a computer readable medium. The examples which follow demonstrate how software which implements the BLAST (Altschul et al., J. Mol. Biol. 215:403-410 (1990)) and BLAZE (Brutlag et al., Comp. Chem. 17:203-207 (1993)) search algorithms on a Sybase system is used to identify open reading frames (ORFs) within a nucleic acid sequence. Such ORFs may be protein encoding fragments and may be useful in producing commercially important proteins such as enzymes used in fermentation reactions and in the production of commercially useful metabolites.

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As used herein, "a computer-based system" refers to the hardware means, software means, and data storage means used to analyze the nucleotide sequence information of the present invention. The minimum hardware means of the computer-based systems of the present invention comprises a central processing unit (CPU), input means, output means, and data storage means. A skilled artisan can readily appreciate that any one of the currently available computer-based systems are suitable for use in the present invention. As stated above, the computer-based systems of the present invention comprise a data storage means having stored therein a nucleotide sequence of the present invention and the necessary hardware means and software means for supporting and implementing a search means. As used herein, "data storage means" refers to memory which can store nucleotide sequence information of the present invention, or a memory access means which can access manufactures having recorded thereon the nucleotide sequence information of the present invention.

As used herein, "search means" refers to one or more programs which are implemented on the computer-based system to compare a target sequence or target structural motif with the sequence information stored within the data storage means. Search means are used to identify fragments or regions of a known sequence which match a particular target sequence or target motif. A variety of known algorithms are disclosed publicly and a variety of commercially available software for conducting search means are and can be used in the computer-based systems of the present invention. Examples of such software includes, but is not limited to, Smith-Waterman, MacPattern (EMBL), BLASTN and BLASTA (NPOLYPEPTIDEIA). A skilled artisan can readily recognize that any one of the available algorithms or implementing software packages for conducting homology searches can be adapted for use in the present computer-based systems. As used herein, a "target sequence" can be any nucleic acid or amino acid sequence of six or more nucleotides or two or more amino acids. A skilled artisan can readily recognize that the longer a target sequence is, the less likely a target sequence will be present as a random occurrence in the database. The most preferred sequence length of a target sequence is from about 10 to 300 amino acids, more preferably from about 30 to 100 nucleotide residues. However, it is well recognized that searches for commercially important fragments,

such as sequence fragments involved in gene expression and protein processing, may be of shorter length.

As used herein, "a target structural motif," or "target motif," refers to any rationally selected sequence or combination of sequences in which the sequence(s) are chosen based on a three-dimensional configuration which is formed upon the folding of the target motif. There are a variety of target motifs known in the art. Protein target motifs include, but are not limited to, enzyme active sites and signal sequences. Nucleic acid target motifs include, but are not limited to, promoter sequences, hairpin structures and inducible expression elements (protein binding sequences).

4.15 TRIPLE HELIX FORMATION

In addition, the fragments of the present invention, as broadly described, can be used to control gene expression through triple helix formation or antisense DNA or RNA, both of which methods are based on the binding of a polynucleotide sequence to DNA or RNA.

Polynucleotides suitable for use in these methods are preferably 20 to 40 bases in length and are designed to be complementary to a region of the gene involved in transcription (triple helix - see Lee et al., Nucl. Acids Res. 6:3073 (1979); Cooney et al., Science 15241:456 (1988); and Dervan et al., Science 251:1360 (1991)) or to the mRNA itself (antisense - Olmno, J. Neurochem. 56:560 (1991); Oligodeoxynucleotides as Antisense Inhibitors of Gene Expression, CRC Press, Boca Raton, FL (1988)). Triple helix-formation optimally results in a shut-off of RNA transcription from DNA, while antisense RNA hybridization blocks translation of an mRNA molecule into polypeptide. Both techniques have been demonstrated to be effective in model systems. Information contained in the sequences of the present invention is necessary for the design of an antisense or triple helix oligonucleotide.

4.16 DIAGNOSTIC ASSAYS AND KITS

The present invention further provides methods to identify the presence or expression of one of the ORFs of the present invention, or homolog thereof, in a test sample, using a nucleic acid probe or antibodies of the present invention, optionally conjugated or otherwise associated with a suitable label.

In general, methods for detecting a polynucleotide of the invention can comprise contacting a sample with a compound that binds to and forms a complex with the polynucleotide for a period sufficient to form the complex, and detecting the complex, so that if a complex is detected, a polynucleotide of the invention is detected in the sample. Such methods can also comprise contacting a sample under stringent hybridization conditions with nucleic acid primers

that anneal to a polynucleotide of the invention under such conditions, and amplifying annealed polynucleotides, so that if a polynucleotide is amplified, a polynucleotide of the invention is detected in the sample.

In general, methods for detecting a polypeptide of the invention can comprise contacting a sample with a compound that binds to and forms a complex with the polypeptide for a period sufficient to form the complex, and detecting the complex, so that if a complex is detected, a polypeptide of the invention is detected in the sample.

In detail, such methods comprise incubating a test sample with one or more of the antibodies or one or more of the nucleic acid probes of the present invention and assaying for binding of the nucleic acid probes or antibodies to components within the test sample.

Conditions for incubating a nucleic acid probe or antibody with a test sample vary. Incubation conditions depend on the format employed in the assay, the detection methods employed, and the type and nature of the nucleic acid probe or antibody used in the assay. One skilled in the art will recognize that any one of the commonly available hybridization, amplification or immunological assay formats can readily be adapted to employ the nucleic acid probes or antibodies of the present invention. Examples of such assays can be found in Chard, T., An Introduction to Radioimmunoassay and Related Techniques, Elsevier Science Publishers. Amsterdam, The Netherlands (1986); Bullock, G.R. et al., Techniques in Immunocytochemistry, Academic Press, Orlando, FL/Vol. 1 (1982), Vol. 2 (1983), Vol. 3 (1985); Tijssen, P., Practice and Theory of immunoassays: Laboratory Techniques in Biochemistry and Molecular Biology, Elsevier Science Publishers, Amsterdam, The Netherlands (1985). The test samples of the present invention include cells, protein or membrane extracts of cells, or biological fluids such as sputum, blood, serum, plasma, or urine. The test sample used in the above-described method will vary based on the assay format, nature of the detection method and the tissues, cells or extracts used as the sample to be assayed. Methods for preparing protein extracts or membrane extracts of cells are well known in the art and can be readily be adapted in order to obtain a sample which is compatible with the system utilized.

In another embodiment of the present invention, kits are provided which contain the necessary reagents to carry out the assays of the present invention. Specifically, the invention provides a compartment kit to receive, in close confinement, one or more containers which comprises: (a) a first container comprising one of the probes or antibodies of the present invention; and (b) one or more other containers comprising one or more of the following: wash reagents, reagents capable of detecting presence of a bound probe or antibody.

In detail, a compartment kit includes any kit in which reagents are contained in separate containers. Such containers include small glass containers, plastic containers or strips of plastic

or paper. Such containers allows one to efficiently transfer reagents from one compartment to another compartment such that the samples and reagents are not cross-contaminated, and the agents or solutions of each container can be added in a quantitative fashion from one compartment to another. Such containers will include a container which will accept the test sample, a container which contains the antibodies used in the assay, containers which contain wash reagents (such as phosphate buffered saline, Tris-buffers, etc.), and containers which contain the reagents used to detect the bound antibody or probe. Types of detection reagents include labeled nucleic acid probes, labeled secondary antibodies, or in the alternative, if the primary antibody is labeled, the enzymatic, or antibody binding reagents which are capable of reacting with the labeled antibody. One skilled in the art will readily recognize that the disclosed probes and antibodies of the present invention can be readily incorporated into one of the established kit formats which are well known in the art.

4.17 MEDICAL IMAGING

The novel polypeptides and binding partners of the invention are useful in medical imaging of sites expressing the molecules of the invention (e.g., where the polypeptide of the invention is involved in the immune response, for imaging sites of inflammation or infection). See, e.g., Kunkel et al., U.S. Pat. NO. 5,413,778. Such methods involve chemical attachment of a labeling or imaging agent, administration of the labeled polypeptide to a subject in a pharmaceutically acceptable carrier, and imaging the labeled polypeptide *in vivo* at the target site.

4.18 SCREENING ASSAYS

Using the isolated proteins and polynucleotides of the invention, the present invention further provides methods of obtaining and identifying agents which bind to a polypeptide encoded by an ORF corresponding to any of the nucleotide sequences set forth in SEQ ID NO: 1-444, or bind to a specific domain of the polypeptide encoded by the nucleic acid. In detail, said method comprises the steps of:

- (a) contacting an agent with an isolated protein encoded by an ORF of the present invention, or nucleic acid of the invention; and
- (b) determining whether the agent binds to said protein or said nucleic acid. In general, therefore, such methods for identifying compounds that bind to a polynucleotide of the invention can comprise contacting a compound with a polynucleotide of the invention for a time sufficient to form a polynucleotide/compound complex, and detecting

the complex, so that if a polynucleotide/compound complex is detected, a compound that binds to a polynucleotide of the invention is identified.

Likewise, in general, therefore, such methods for identifying compounds that bind to a polypeptide of the invention can comprise contacting a compound with a polypeptide of the invention for a time sufficient to form a polypeptide/compound complex, and detecting the complex, so that if a polypeptide/compound complex is detected, a compound that binds to a polynucleotide of the invention is identified.

Methods for identifying compounds that bind to a polypeptide of the invention can also comprise contacting a compound with a polypeptide of the invention in a cell for a time sufficient to form a polypeptide/compound complex, wherein the complex drives expression of a receptor gene sequence in the cell, and detecting the complex by detecting reporter gene sequence expression, so that if a polypeptide/compound complex is detected, a compound that binds a polypeptide of the invention is identified.

Compounds identified via such methods can include compounds which modulate the activity of a polypeptide of the invention (that is, increase or decrease its activity, relative to activity observed in the absence of the compound). Alternatively, compounds identified via such methods can include compounds which modulate the expression of a polynucleotide of the invention (that is, increase or decrease expression relative to expression levels observed in the absence of the compound). Compounds, such as compounds identified via the methods of the invention, can be tested using standard assays well known to those of skill in the art for their ability to modulate activity/expression.

The agents screened in the above assay can be, but are not limited to, peptides, carbohydrates, vitamin derivatives, or other pharmaceutical agents. The agents can be selected and screened at random or rationally selected or designed using protein modeling techniques.

For random screening, agents such as peptides, carbohydrates, pharmaceutical agents and the like are selected at random and are assayed for their ability to bind to the protein encoded by the ORF of the present invention. Alternatively, agents may be rationally selected or designed. As used herein, an agent is said to be "rationally selected or designed" when the agent is chosen based on the configuration of the particular protein. For example, one skilled in the art can readily adapt currently available procedures to generate peptides, pharmaceutical agents and the like, capable of binding to a specific peptide sequence, in order to generate rationally designed antipeptide peptides, for example see Hurby et al., Application of Synthetic Peptides: Antisense Peptides," In Synthetic Peptides, A User's Guide, W.H. Freeman, NY (1992), pp. 289-307, and Kaspczak et al., Biochemistry 28:9230-8 (1989), or pharmaceutical agents, or the like.

In addition to the foregoing, one class of agents of the present invention, as broadly described, can be used to control gene expression through binding to one of the ORFs or EMFs of the present invention. As described above, such agents can be randomly screened or rationally designed/selected. Targeting the ORF or EMF allows a skilled artisan to design sequence specific or element specific agents, modulating the expression of either a single ORF or multiple ORFs which rely on the same EMF for expression control. One class of DNA binding agents are agents which contain base residues which hybridize or form a triple helix formation by binding to DNA or RNA. Such agents can be based on the classic phosphodiester, ribonucleic acid backbone, or can be a variety of sulfhydryl or polymeric derivatives which have base attachment capacity.

Agents suitable for use in these methods preferably contain 20 to 40 bases and are designed to be complementary to a region of the gene involved in transcription (triple helix - see Lee et al., Nucl. Acids Res. 6:3073 (1979); Cooney et al., Science 241:456 (1988); and Dervan et al., Science 251:1360 (1991)) or to the mRNA itself (antisense - Okano, J. Neurochem. 56:560 (1991); Oligodeoxynucleotides as Antisense Inhibitors of Gene Expression, CRC Press, Boca Raton, FL (1988)). Triple helix-formation optimally results in a shut-off of RNA transcription from DNA, while antisense RNA hybridization blocks translation of an mRNA molecule into polypeptide. Both techniques have been demonstrated to be effective in model systems. Information contained in the sequences of the present invention is necessary for the design of an antisense or triple helix oligonucleotide and other DNA binding agents.

Agents which bind to a protein encoded by one of the ORFs of the present invention can be used as a diagnostic agent. Agents which bind to a protein encoded by one of the ORFs of the present invention can be formulated using known techniques to generate a pharmaceutical composition.

4.19 USE OF NUCLEIC ACIDS AS PROBES

Another aspect of the subject invention is to provide for polypeptide-specific nucleic acid hybridization probes capable of hybridizing with naturally occurring nucleotide sequences. The hybridization probes of the subject invention may be derived from any of the nucleotide sequences SEQ ID NO: 1-444. Because the corresponding gene is only expressed in a limited number of tissues, a hybridization probe derived from any of the nucleotide sequences SEQ ID NO: 1-444 can be used as an indicator of the presence of RNA of cell type of such a tissue in a sample.

Any suitable hybridization technique can be employed, such as, for example, in situ hybridization. PCR as described in US Patents Nos. 4,683,195 and 4,965,188 provides

additional uses for oligonucleotides based upon the nucleotide sequences. Such probes used in PCR may be of recombinant origin, may be chemically synthesized, or a mixture of both. The probe will comprise a discrete nucleotide sequence for the detection of identical sequences or a degenerate pool of possible sequences for identification of closely related genomic sequences.

Other means for producing specific hybridization probes for nucleic acids include the cloning of nucleic acid sequences into vectors for the production of mRNA probes. Such vectors are known in the art and are commercially available and may be used to synthesize RNA probes in vitro by means of the addition of the appropriate RNA polymerase as T7 or SP6 RNA polymerase and the appropriate radioactively labeled nucleotides. The nucleotide sequences may be used to construct hybridization probes for mapping their respective genomic sequences. The nucleotide sequence provided herein may be mapped to a chromosome or specific regions of a chromosome using well known genetic and/or chromosomal mapping techniques. These techniques include in situ hybridization, linkage analysis against known chromosomal markers, hybridization screening with libraries or flow-sorted chromosomal preparations specific to known chromosomes, and the like. The technique of fluorescent in situ hybridization of chromosome spreads has been described, among other places, in Verma et al (1988) Human Chromosomes: A Manual of Basic Techniques, Pergamon Press, New York NY.

Fluorescent in situ hybridization of chromosomal preparations and other physical chromosome mapping techniques may be correlated with additional genetic map data. Examples of genetic map data can be found in the 1994 Genome Issue of Science (265:1981f). Correlation between the location of a nucleic acid on a physical chromosomal map and a specific disease (or predisposition to a specific disease) may help delimit the region of DNA associated with that genetic disease. The nucleotide sequences of the subject invention may be used to detect differences in gene sequences between normal, carrier or affected individuals.

4.20 PREPARATION OF SUPPORT BOUND OLIGONUCLEOTIDES

Oligonucleotides, i.e., small nucleic acid segments, may be readily prepared by, for example, directly synthesizing the oligonucleotide by chemical means, as is commonly practiced using an automated oligonucleotide synthesizer.

Support bound oligonucleotides may be prepared by any of the methods known to those of skill in the art using any suitable support such as glass, polystyrene or Teflon. One strategy is to precisely spot oligonucleotides synthesized by standard synthesizers. Immobilization can be achieved using passive adsorption (Inouye & Hondo, (1990) J. Clin. Microbiol. 28(6) 1469-72); using UV light (Nagata et al., 1985; Dahlen et al., 1987; Morrissey & Collins, (1989) Mol. Cell

Probes 3(2) 189-207) or by covalent binding of base modified DNA (Keller et al., 1988; 1989); all references being specifically incorporated herein.

Another strategy that may be employed is the use of the strong biotin-streptavidin interaction as a linker. For example, Broude et al. (1994) Proc. Natl. Acad. Sci. USA 91(8) 3072-6, describe the use of biotinylated probes, although these are duplex probes, that are immobilized on streptavidin-coated magnetic beads. Streptavidin-coated beads may be purchased from Dynal, Oslo. Of course, this same linking chemistry is applicable to coating any surface with streptavidin. Biotinylated probes may be purchased from various sources, such as, e.g., Operon Technologies (Alameda, CA).

Nunc Laboratories (Naperville, IL) is also selling suitable material that could be used. Nunc Laboratories have developed a method by which DNA can be covalently bound to the microwell surface termed Covalink NH. CovaLink NH is a polystyrene surface grafted with secondary amino groups (>NH) that serve as bridge-heads for further covalent coupling. CovaLink Modules may be purchased from Nunc Laboratories. DNA molecules may be bound to CovaLink exclusively at the 5'-end by a phosphoramidate bond, allowing immobilization of more than 1 pmol of DNA (Rasmussen et al., (1991) Anal. Biochem. 198(1) 138-42).

The use of CovaLink NH strips for covalent binding of DNA molecules at the 5'-end has been described (Rasmussen et al., (1991). In this technology, a phosphoramidate bond is employed (Chu et al., (1983) Nucleic Acids Res. 11(8) 6513-29). This is beneficial as immobilization using only a single covalent bond is preferred. The phosphoramidate bond joins the DNA to the CovaLink NH secondary amino groups that are positioned at the end of spacer arms covalently grafted onto the polystyrene surface through a 2 nm long spacer arm. To link an oligonucleotide to CovaLink NH via an phosphoramidate bond, the oligonucleotide terminus must have a 5'-end phosphate group. It is, perhaps, even possible for biotin to be covalently bound to CovaLink and then streptavidin used to bind the probes.

More specifically, the linkage method includes dissolving DNA in water (7.5 ng/µl) and denaturing for 10 min. at 95°C and cooling on ice for 10 min. Ice-cold 0.1 M 1-methylimidazole, pH 7.0 (1-MeIm₇), is then added to a final concentration of 10 mM 1-MeIm₇. The single-stranded DNA solution is then dispensed into CovaLink NH strips (75 µl/well) standing on ice.

Carbodiimide 0.2 M 1-ethyl-3-(3-dimethylaminopropyl)-carbodiimide (EDC), dissolved in 10 mM 1-MeIm7, is made fresh and 25 µl added per well. The strips are incubated for 5 hours at 50°C. After incubation the strips are washed using, e.g., Nunc-Immuno Wash; first the wells are washed 3 times, then they are soaked with washing solution for 5 min., and finally they are washed 3 times (where in the washing solution is 0.4 N NaOH, 0.25% SDS heated to 50°C).

It is contemplated that a further suitable method for use with the present invention is that described in PCT Patent Application WO 90/03382 (Southern & Maskos), incorporated herein by reference. This method of preparing an oligonucleotide bound to a support involves attaching a nucleoside 3'-reagent through the phosphate group by a covalent phosphodiester link to aliphatic hydroxyl groups carried by the support. The oligonucleotide is then synthesized on the supported nucleoside and protecting groups removed from the synthetic oligonucleotide chain under standard conditions that do not cleave the oligonucleotide from the support. Suitable reagents include nucleoside phosphoramidite and nucleoside hydrogen phosphorate.

An on-chip strategy for the preparation of DNA probe for the preparation of DNA probe arrays may be employed. For example, addressable laser-activated photodeprotection may be employed in the chemical synthesis of oligonucleotides directly on a glass surface, as described by Fodor *et al.* (1991) Science 251(4995) 767-73, incorporated herein by reference. Probes may also be immobilized on nylon supports as described by Van Ness *et al.* (1991) Nucleic Acids Res. 19(12) 3345-50; or linked to Teflon using the method of Duncan & Cavalier (1988) Anal. Biochem. 169(1) 104-8; all references being specifically incorporated herein.

To link an oligonucleotide to a nylon support, as described by Van Ness *et al.* (1991), requires activation of the nylon surface via alkylation and selective activation of the 5'-amine of oligonucleotides with cyanuric chloride.

One particular way to prepare support bound oligonucleotides is to utilize the light-generated synthesis described by Pease *et al.*, (1994) PNAS USA 91(11) 5022-6, incorporated herein by reference). These authors used current photolithographic techniques to generate arrays of immobilized oligonucleotide probes (DNA chips). These methods, in which light is used to direct the synthesis of oligonucleotide probes in high-density, miniaturized arrays, utilize photolabile 5'-protected *N*-acyl-deoxynucleoside phosphoramidites, surface linker chemistry and versatile combinatorial synthesis strategies. A matrix of 256 spatially defined oligonucleotide probes may be generated in this manner.

4.21 PREPARATION OF NUCLEIC ACID FRAGMENTS

The nucleic acids may be obtained from any appropriate source, such as cDNAs, genomic DNA, chromosomal DNA, microdissected chromosome bands, cosmid or YAC inserts, and RNA, including mRNA without any amplification steps. For example, Sambrook *et al.* (1989) describes three protocols for the isolation of high molecular weight DNA from mammalian cells (p. 9.14-9.23).

DNA fragments may be prepared as clones in M13, plasmid or lambda vectors and/or prepared directly from genomic DNA or cDNA by PCR or other amplification methods. Samples

may be prepared or dispensed in multiwell plates. About 100-1000 ng of DNA samples may be prepared in 2-500 ml of final volume.

The nucleic acids would then be fragmented by any of the methods known to those of skill in the art including, for example, using restriction enzymes as described at 9.24-9.28 of Sambrook *et al.* (1989), shearing by ultrasound and NaOH treatment.

Low pressure shearing is also appropriate, as described by Schriefer *et al.* (1990) Nucleic Acids Res. 18(24) 7455-6, incorporated herein by reference). In this method, DNA samples are passed through a small French pressure cell at a variety of low to intermediate pressures. A lever device allows controlled application of low to intermediate pressures to the cell. The results of these studies indicate that low-pressure shearing is a useful alternative to sonic and enzymatic DNA fragmentation methods.

One particularly suitable way for fragmenting DNA is contemplated to be that using the two base recognition endonuclease, *CviII*, described by Fitzgerald *et al.* (1992) Nucleic Acids Res. 20(14) 3753-62. These authors described an approach for the rapid fragmentation and fractionation of DNA into particular sizes that they contemplated to be suitable for shotgun cloning and sequencing.

The restriction endonuclease CviJI normally cleaves the recognition sequence PuGCPy between the G and C to leave blunt ends. Atypical reaction conditions, which alter the specificity of this enzyme (CviJI**), yield a quasi-random distribution of DNA fragments form the small molecule pUC19 (2688 base pairs). Fitzgerald et al. (1992) quantitatively evaluated the randomness of this fragmentation strategy, using a CviJI** digest of pUC19 that was size fractionated by a rapid gel filtration method and directly ligated, without end repair, to a lac Z minus M13 cloning vector. Sequence analysis of 76 clones showed that CviJI** restricts pyGCPy and PuGCPu, in addition to PuGCPy sites, and that new sequence data is accumulated at a rate consistent with random fragmentation.

As reported in the literature, advantages of this approach compared to sonication and agarose gel fractionation include: smaller amounts of DNA are required (0.2-0.5 μ g instead of 2-5 μ g); and fewer steps are involved (no preligation, end repair, chemical extraction, or agarose gel electrophoresis and elution are needed

Irrespective of the manner in which the nucleic acid fragments are obtained or prepared, it is important to denature the DNA to give single stranded pieces available for hybridization. This is achieved by incubating the DNA solution for 2-5 minutes at 80-90°C. The solution is then cooled quickly to 2°C to prevent renaturation of the DNA fragments before they are contacted with the chip. Phosphate groups must also be removed from genomic DNA by methods known in the art.

4.22 PREPARATION OF DNA ARRAYS

Arrays may be prepared by spotting DNA samples on a support such as a nylon membrane. Spotting may be performed by using arrays of metal pins (the positions of which correspond to an array of wells in a microtiter plate) to repeated by transfer of about 20 nl of a DNA solution to a nylon membrane. By offset printing, a density of dots higher than the density of the wells is achieved. One to 25 dots may be accommodated in 1 mm², depending on the type of label used. By avoiding spotting in some preselected number of rows and columns, separate subsets (subarrays) may be formed. Samples in one subarray may be the same genomic segment of DNA (or the same gene) from different individuals, or may be different, overlapped genomic clones. Each of the subarrays may represent replica spotting of the same samples. In one example, a selected gene segment may be amplified from 64 patients. For each patient, the amplified gene segment may be in one 96-well plate (all 96 wells containing the same sample). A plate for each of the 64 patients is prepared. By using a 96-pin device, all samples may be spotted on one 8 x 12 cm membrane. Subarrays may contain 64 samples, one from each patient. Where the 96 subarrays are identical, the dot span may be 1 mm² and there may be a 1 mm space between subarrays.

Another approach is to use membranes or plates (available from NUNC, Naperville, Illinois) which may be partitioned by physical spacers e.g. a plastic grid molded over the membrane, the grid being similar to the sort of membrane applied to the bottom of multiwell plates, or hydrophobic strips. A fixed physical spacer is not preferred for imaging by exposure to flat phosphor-storage screens or x-ray films.

The present invention is illustrated in the following examples. Upon consideration of the present disclosure, one of skill in the art will appreciate that many other embodiments and variations may be made in the scope of the present invention. Accordingly, it is intended that the broader aspects of the present invention not be limited to the disclosure of the following examples. The present invention is not to be limited in scope by the exemplified embodiments which are intended as illustrations of single aspects of the invention, and compositions and methods which are functionally equivalent are within the scope of the invention. Indeed, numerous modifications and variations in the practice of the invention are expected to occur to those skilled in the art upon consideration of the present preferred embodiments. Consequently, the only limitations which should be placed upon the scope of the invention are those which appear in the appended claims.

All references cited within the body of the instant specification are hereby incorporated by reference in their entirety.

5. EXAMPLES

5.1 EXAMPLE 1

Novel Nucleic Acid Sequences Obtained From Various Libraries

A plurality of novel nucleic acids were obtained from cDNA libraries prepared from various human tissues and in some cases isolated from a genomic library derived from human chromosome using standard PCR, SBH sequence signature analysis and Sanger sequencing techniques. The inserts of the library were amplified with PCR using primers specific for the vector sequences which flank the inserts. Clones from cDNA libraries were spotted on nylon membrane filters and screened with oligonucleotide probes (e.g., 7-mers) to obtain signature sequences. The clones were clustered into groups of similar or identical sequences. Representative clones were selected for sequencing.

In some cases, the 5' sequence of the amplified inserts was then deduced using a typical Sanger sequencing protocol. PCR products were purified and subjected to fluorescent dye terminator cycle sequencing. Single pass gel sequencing was done using a 377 Applied Biosystems (ABI) sequencer to obtain the novel nucleic acid sequences

5.2 EXAMPLE 2

Assemblage of Novel Nucleic Acids

The nucleic acids of the present invention, designated as SEQ ID NO: 1-444 were assembled using an EST sequence as a seed. Then a recursive algorithm was used to extend the seed EST into an extended assemblage, by pulling additional sequences from different databases (i.e., Hyseq's database containing EST sequences, dbEST, gb pri, and UniGene, and exons from public domain genomic sequences predicated by GenScan) that belong to this assemblage. The algorithm terminated when there was no additional sequences from the above databases that would extend the assemblage. Further, inclusion of component sequences into the assemblage was based on a BLASTN hit to the extending assemblage with BLAST score greater than 300 and percent identity greater than 95%.

Using PHRAP (Univ. of Washington) or CAP4 (Paracel), full-length gene sequences and their corresponding protein sequences were generated from the assemblage. Any frame shifts and incorrect stop codons were corrected by hand editing. During editing, the sequence was checked using FASTXY algorithm against Genbank (i.e., dbEST, gb pri, UniGene, and Genpept). Other computer programs which may have been used in the editing process were phredPhrap and Consed (University of Washington) and ed-ready, ed-ext and gc-zip-2 (Hyseq, Inc.). The full-length nucleotide sequences are shown in the Sequence Listing as SEQ ID NO: 1-444. The corresponding polypeptide sequences are SEQ ID NO: 445-888.

Table 1 shows the various tissue sources of SEQ ID NO: 1-444.

The nearest neighbor results for polypeptides encoded by SEQ ID NO: 1-444 (i.e. SEQ ID NO: 445-888) were obtained by a BLASTP (version 2.0al 19MP-WashU) search against Genpept release 124 using BLAST algorithm. The nearest neighbor result showed the closest homologue with functional annotation for SEQ ID NO: 1-444 from Genpept. The translated amino acid sequences for which the nucleic acid sequence encodes are shown in the Sequence Listing. The homologues with identifiable functions for SEQ ID NO: 1-444 are shown in Table 2 below.

Using eMatrix software package (Stanford University, Stanford, CA) (Wu et al., J. Comp. Biol., Vol. 6 pp. 219-235 (1999) herein incorporated by reference), polypeptides encoded by SEQ ID NO: 1-444 (i.e. SEQ ID NO: 445-888) were examined to determine whether they had identifiable signature regions. Table 3 shows the signature region found in the indicated polypeptide sequences, the description of the signature, the eMatrix p-value(s) and the position(s) of the signature within the polypeptide sequence.

Using the pFam software program (Sonnhammer et al., Nucleic Acids Res., Vol. 26(1) pp. 320-322 (1998) herein incorporated by reference) polypeptides encoded by SEQ ID NO: 1-444 (i.e. SEQ ID NO: 445-888) were examined for domains with homology to certain peptide domains. Table 4 shows the name of the domain found, the description, the p-value and the pFam score for the identified domain within the sequence.

The GeneAtlas™ software package (Molecular Simulations Inc. (MSI), San Diego, CA) was used to predict the three-dimensional structure models for the polypeptides encoded by SEQ ID NO: 1-444 (i.e. SEQ ID NO: 445-888). Models were generated by (1) PSI-BLAST which is a multiple alignment sequence profile-based searching developed by Altschul et al, (Nucl. Acids. Res. 25, 3389-3408 (1997)), (2) High Throughput Modeling (HTM) (Molecular Simulations Inc. (MSI) San Diego, CA,) which is an automated sequence and structure searching procedure (http://www.msi.com/), and (3) SeqFold™ which is a fold recognition method described by Fischer and Eisenberg (J. Mol. Biol. 209, 779-791 (1998)). This analysis was carried out, in part, by comparing the polypeptides of the invention with the known NMR (nuclear magnetic resonance) and x-ray crystal three-dimensional structures as templates. Table 5 shows, "PDB ID", the Protein DataBase (PDB) identifier given to template structure; "Chain ID", identifier of the subcomponent of the PDB template structure; "Compound Information", information of the PDB template structure and/or its subcomponents; "PDB Function Annotation" gives function of the PDB template as annotated by the PDB files (http://www.rcsb.org/PDB/); start and end amino acid position of the protein sequence aligned; PSI-BLAST score, the verify score, the SeqFold score, and the Potential(s) of Mean Force (PMF). The verify score is produced by GeneAtlas™

software (MSI), is based on Dr. Eisenberg's Profile-3D threading program developed in Dr. David Eisenberg's laboratory (US patent no. 5,436,850 and Luthy, Bowie, and Eisenberg, Nature, 356:83-85 (1992)) and a publication by R. Sanchez and A. Sali, Proc. Natl. Acad. Sci. USA, 95:13597-12502. The verify score produced by GeneAtlas normalizes the verify score for proteins with different lengths so that a unified cutoff can be used to select good models as follows:

Verify score (normalized) = (raw score - 1/2 high score)/(1/2 high score)

The PFM score, produced by GeneAtlas™ software (MSI), is a composite scoring function that depends in part on the compactness of the model, sequence identity in the alignment used to build the model, pairwise and surface mean force potentials (MFP). As given in table 8, a verify score between 0 to 1.0, with 1 being the best, represents a good model. Similarly, a PMF score between 0 to 1.0, with 1 being the best, represents a good model. A SeqFold™ score of more than 50 is considered significant. A good model may also be determined by one of skill in the art based all the information in Table 5 taken in totality.

The nucleotide sequence within the sequences that codes for signal peptide sequences and their cleavage sites can be determined from using Neural Network SignalP V1.1 program (from Center for Biological Sequence Analysis, The Technical University of Denmark). The process for identifying prokaryotic and eukaryotic signal peptides and their cleavage sites are also disclosed by Henrik Nielson, Jacob Engelbrecht, Soren Brunak, and Gunnar von Heijne in the publication "Identification of prokaryotic and eukaryotic signal peptides and prediction of their cleavage sites" Protein Engineering, Vol. 10, no. 1, pp. 1-6 (1997), incorporated herein by reference. A maximum S score and a mean S score, as described in the Nielson et al, as reference, were obtained for the polypeptide sequences. Table 6 shows the position of the last amino acid of the signal peptide in each of the polypeptides and the maximum score and mean score associated with that signal peptide.

Table 7 correlates each of SEQ ID NO: 1-444 to a specific chromosomal location.

Table 8 is a correlation table of the novel polynucleotide sequences SEQ ID NO: 1-444, and their corresponding priority nucleotide sequences in the priority application USSN 09/659,671, herein incorporated by reference in its entirety.

TABLE 1

Tissue Origin	RNA Source	Library Name	SEQ ID NO:
adult brain	GIBCO	AB3001	4 6-8 12 23 33-34 47 50 55 57-60 62 89 102 104-106 123 144 162 176-177 179 187 194 248 260 270 279 292 294 297-298 307 322-323 326 333 336 341 351 450
adult brain	GIBCO	ABD003	6 10 12-15 17-18 26 31 34-35 38-40 42-44 46 48-50 53 56 59-60 64 66 70-72 80-81 85-86 98 101 107 116-117 125 130 138-139 142 144 147 151 160-161 164 173 175-177 179 184-185 187-188 194-195 198 201 215 217-218 222 226 228 232 239-240 243-244 247 252 256 258 260 264-265 267 274-275 284 288 290 293 298 306-308 314-315 318-320 325-326 333-334 337 341 343 345-346 351-354 364-365 371 390-391 424-425 429
adult brain	Clontech	ABR001	5 36 43 76 108 128 182-183 212 239 242 260 263 269 296 325 351 364 371-372 423
adult brain	Clontech	ABR006	2 9 11 13 18 23 35 38 42 46 50-51 53-54 60 63 66 85 91 107-108 116-117 120 122 128 170 178 180 184 187-188 193-194 198 202 215 232 243 245 257-258 260 266-267 271 285 294 301 333-334 337 370 389 394-396 400 405 412 423 428 434 436 453 458
adult brain	Clontech	ABR008	1 3 7 10-14 16-17 19-23 26-28 34-35 38-39 41 43 46-48 51-54 56 60 62 64 66-68 75 82 86-87 91 96-98 102 104-106 108 110-111 114 116-118 122 125 127-130 134 138-139 141-143 145-146 150-151 153 156 158 160 162 167-170 173-175 177-180 182 185-186 191-194 196-197 200-201 205-206 208-209 211 213-215 219-220 226-227 231 238 241 244 246-248 252 256 260 262-265 269 271 273 278-280 282 284 290 292 296 298 301-302 306 309 311 315-317 322-323 325-327 329-331 335-337 339 342-343 345-346 350-355 359-360 362 364 368 370 372 374 376 381 383 385 387 390-395 400-401 405 410 412 414 417 420-421 423-425 432 440 447 450-452 459 472-473
adult brain	Clontech	ABR011	174 177 360
adult brain	BioChain	ABR012	334 341
adult brain	BioChain	ABR013	41-42 60 101 163 355
adult brain adult brain	Invitrogen Invitrogen	ABR014 ABR015	53 95 104-106 143 149 177 180 258 42 70-72 79 95 112 138-140 163 195 275 288 322-323 341 343 458
adult brain	Invitrogen	ABR016	13 31 60 79 124 136 154 163 333 341 343 364 370
adult brain	Invitrogen	ABT004	1 11-13 15 18 24-26 34 50 56 68 87 98 104- 106 111 123-124 131-133 137 144 146 173 189 194 206 224 247-248 260 262 264 269 272 274 282 298 318 327 335 346 351 356- 357 372 375 381 392 409-410 421
cultured preadipocytes	Stratagene	ADP001	2 11-14 24-25 40 42 47 50 52 57-58 69 76 107 120 144 151 156 163 168 171 194 197 199 203-204 215 229 250-251 262 294 333 338 341 415 450 469-473
adrenal gland	Clontech	ADR002	10-11 16 18 22-23 27-28 33-35 40 43-45 49 61 66 85 98 107-108 111 116-117 124 136 143 145 160 167 173 175 184 187 201 217-

			218 229 249-251 258 262 269 271 273 277
]	280 287 289 298 301 308 322-323 337 352
			354 360 414 425 445-446 463
adult heart	GIBCO ,	AHR001	11-13 15 20-23 26-27 30 33-34 37-40 49 53
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			125 129 131-145 147 151 154 156 162-165
	1	1	172-173 175-180 182-191 193-195 197-206
	1		208-217 223 227 233 239-240 242-244 247
		1	258-259 261 264 267 269 273 277 282 286
	1		288 290 301-302 306 310 313 325 333-334
	İ		336-337 341 343 345 351 364 369-372 469-
	1		473
macrophage	Invitrogen	HMP001	49 123 144 151 275
macrophage	Invitrogen		
infant brain	Columbia	IB2002	7-8 11-13 16 20-22 24-26 34 38 44-58 60
	University		62-64 66 68 75 80 84 87 91-92 94 97-98
			101 103-108 123 126 128 130-133 135-137
	.1	l	142-146 148 151 156 158 160 164 170 173

			175 177-178 180 182 187-188 194 196-198
			200-201 206 212 215 217 219 226 232-233
			239 241 247-248 256 260 263 265 268 273
			277-278 282-284 286 288-289 294 298
			301-303 306-307 309 312 324 330 334-335
			337-339 342-343 346-347 351 355 364 370
			373-376 389-394 400 413 421 423-424 458
			469-471
infant brain	Columbia	IB2003	4 8 11 13-14 16 23 42 46-47 50 54 56-62
	University		65-66 76 94 98 102 119-120 135 142-143
			145 150-152 158 163-164 173 175 180 226
			233 244 247 260 262 267 277 302 304 309
			319-320 334-337 351 364 375 383 389-392
			400 423 427 434 472-473
infant brain	Columbia	1BM002	33 50 54 112 131-133 163 173 215 226 267
	University		331 423
infant brain	Columbia	IBS001	2 5 11 26 34 52 87 91 98 108 170 173 177
	University	<u> </u>	194 200-201 248 277 340 361 412 423
lung, fibroblast	Stratagene	LFB001	13 16-17 22 26 39 46 57-58 78 83 88 93
	٠.		101 116 122 131-133 160 170 178 195 198
	į,		210 214 223 262 267 276 304 319-320 322-
			323 333 341 349 375 383 417 447 455
lung tumor	Invitrogen	LGT002	10-12 15 17-18 20 23 26-28 30 32 34 37
	i		39-40 43 49 51 56-58 62 64 66 80 85-87 91
	1		94 98 101-102 104-106 108 111 122 124
			126-129 134 136 142-144 147 156-157 168
	[ļ	173-176 179 184 186-187 189 195 197-198
•		1	203-204 209-210 218 220-222 226 232-233
	Ĭ		237-239 244-246 249-251 253 257 259 269
		}	273 277-279 282-284 287 300-301 308 310
		1	314 319-320 327 333-335 341 346 348 352
	1	1	358 362 369 371 377-379 392 394 397 406-
	·	 	407 410 412 421-422 436 469-471
lymphocytes	ATCC	LPC001	7 10-11 14-15 18 20 24-25 27 33 35 43-44
			49 57-58 65-66 74-75 80 88 95 103-106
	ì	i	108 113 116 124 130-133 145-146 151 163-
			164 167-171 185 200 206 215 218 226 228
		1	232 241 244 247-248 262 267 273 275-277
			284 297 321 331 345 349 352 375 400 472-
ļ.,	01000	7710001	473
leukocyte	GIBCO ·	LUC001	1 6-8 11 14-15 17-18 20 23-25 27-35 39 43
			46 49 52-54 57-58 62-63 74 76-78 80-81
	1	Į.	84-85 88 90 92-95 98 102 104-106 108 112
ł	ļ	1	114 117 119 123-125 128 130-135 141 143-
	}		146 148 151 153 156-158 160-164 166-168
	Ì		171 174-176 178-179 181 183-184 187-188
			195 199-200 202-204 206 209 211-216
			218-219 221-223 226-228 232-235 239-240
	İ		242 244-245 247 256 258 260-262 264 266-
	1	ļ	267 270 275-277 279-280 282 284-290
1			292-293 297-298 300-302 306 308-310 312
l .			314 317-328 330 333 335 341 346 349 400
loukeaute	Clastach	T LICOO2	412 427 436-437 450 462 469-473 17 19 27 34 42-43 46 49 90 98 104-106 108
leukocyte	Clontech	LUC003	•
	1		113 122-123 128 157 206 284-285 321 333
Malanana San	Clastes	NOT DOA	341 362 472-473 6 11 30 34 45 54-55 61-62 65-66 78 81 93
Melanoma from	Clontech	MEL004	6 11 30 34 45 54-55 61-62 65-66 78 81 93
cell line ATCC]	ļ	112 114 116 122 128 130 135 143 145 160 164 177 180 187 195 219 227 235 239-240
#CRL 1424	Ĭ		1
			258 264-265 279-280 287 302 304 306 311
	Invitrogen	MMG001	1

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		Ţ	40 43 45 47 49-51 53-54 57-58 62 64 66
		İ	70-72 76 80 83 87-88 95-96 98-99 101-102
			104-106 108-109 112 118 124-125 127-133
	İ	ì	136 138-139 142-146 148 150-151 163 167
	Í	ļ	170-171 175 180 186-187 189 197 200 212
		\	224 226-227 241 244 247-248 250-251
			253-255 257-258 260 262-263 265 269 272
			276-280 282-283 288 290 301 306-307 309
			313 315-316 322-323 326-327 333-334 337
			341 343-346 356 361 364 370 387 390-392
			404 409 412 415 421 428 430-432 469-473
induced neuron	Stratagene	NTD001	7 20 42 47 49 53 83 121 134 136 151 153
cells			195 202 218-219 223 247 264 267 269 302
			312-313 339 382 427 444
retinoic acid-	Stratagene	NTR001	34 70-72 104-106 110 116 197 258 392 396
induced neuronal			422
cells	<u> </u>		·
neuronal cells	Stratagene	NTU001	16 40 49 53-54 80 100 130 136 194 258
		<u> </u>	281 396 427 472-473
pituitary gland	Clontech	PIT004	54 119 170 200 242 264 270 319-320 333
			336 341 414
placenta	Clontech	PLA003	32 304 341 421
prostate	Clontech	PRT001	9 13 21 23 53 59 79 83 88 119 123 127 160
			162 178 180 182-183 187 209 250-251 273
			278 292-293 306 329 333 336 409 442
rectum	Invitrogen	REC001	5 12 15 22 32 42 80 108 118 127 143-144
		Į	187 194 224 226 277 298 345 396 404 442
	1		444
salivary gland	Clontech	SAL001	4 7 10 40 66 88 102 104-106 126 128 151
			162 182 212 222 242 252 277 287 312 319-
			320 348 369
salivary gland	Clontech	SALS03	42
skin fibroblast	ATCC	SFB001	54
skin fibroblast	ATCC	SFB003	87 144
small intestine	Clontech	SIN001	1 9 11 13 15 17-18 22 27 33 38 40 54 57-58
		1	61 63-64 78-80 84 97 102 115-116 124
		1	128-129 131-134 142 144 148 153 159-160
1		į	163 165 167 180 186 195 197-198 202-204
	1	1	214-215 217 230 232 234 242-243 248 257
		1	265 271 278 284 295 297 299 304 306 310-
		1	311 319-321 328 333 337 341 358 372-373
	61		395 408 438 461 467
skeletal muscle			12 64 76 07 00 104 106 160 176 000 000
0	Clontech	SKM001	13 64 76 87 92 104-106 122 176 202 302
*			322-323 327 341 451 464 ·
skeletal muscle	Clontech	SKM002	322-323 327 341 451 464 246
skeletal muscle skeletal muscle	Clontech Clontech	SKM002 SKMS03	322-323 327 341 451 464 246 49
skeletal muscle skeletal muscle skeletal muscle	Clontech Clontech null	SKM002 SKMS03 SKMS04	322-323 327 341 451 464 246 49 88
skeletal muscle skeletal muscle	Clontech Clontech	SKM002 SKMS03	322-323 327 341 451 464 246 49 88 6 12 22-23 48 60 70-72 80 93 101-102 104-
skeletal muscle skeletal muscle skeletal muscle	Clontech Clontech null	SKM002 SKMS03 SKMS04	322-323 327 341 451 464 246 49 88 6 12 22-23 48 60 70-72 80 93 101-102 104- 106 114 125 135 138-139 143-144 148 162-
skeletal muscle skeletal muscle skeletal muscle	Clontech Clontech null	SKM002 SKMS03 SKMS04	322-323 327 341 451 464 246 49 88 6 12 22-23 48 60 70-72 80 93 101-102 104- 106 114 125 135 138-139 143-144 148 162- 163 167 176 196 200 202-204 242-243 264
skeletal muscle skeletal muscle skeletal muscle	Clontech Clontech null	SKM002 SKMS03 SKMS04	322-323 327 341 451 464 246 49 88 6 12 22-23 48 60 70-72 80 93 101-102 104- 106 114 125 135 138-139 143-144 148 162- 163 167 176 196 200 202-204 242-243 264 270 283 288 294 302 321 326 329 333 336
skeletal muscle skeletal muscle skeletal muscle spinal cord	Clontech Clontech null Clontech	SKM002 SKMS03 SKMS04 SPC001	322-323 327 341 451 464 246 49 88 6 12 22-23 48 60 70-72 80 93 101-102 104- 106 114 125 135 138-139 143-144 148 162- 163 167 176 196 200 202-204 242-243 264 270 283 288 294 302 321 326 329 333 336 343 345 349 352 354 382 424-425 436 457
skeletal muscle skeletal muscle skeletal muscle spinal cord adult spleen	Clontech Clontech null Clontech	SKM002 SKMS03 SKMS04 SPC001	322-323 327 341 451 464 246 49 88 6 12 22-23 48 60 70-72 80 93 101-102 104- 106 114 125 135 138-139 143-144 148 162- 163 167 176 196 200 202-204 242-243 264 270 283 288 294 302 321 326 329 333 336 343 345 349 352 354 382 424-425 436 457 2 98 120 141 164 166 244 260 278 394
skeletal muscle skeletal muscle skeletal muscle spinal cord	Clontech Clontech null Clontech	SKM002 SKMS03 SKMS04 SPC001	322-323 327 341 451 464 246 49 88 6 12 22-23 48 60 70-72 80 93 101-102 104- 106 114 125 135 138-139 143-144 148 162- 163 167 176 196 200 202-204 242-243 264 270 283 288 294 302 321 326 329 333 336 343 345 349 352 354 382 424-425 436 457 2 98 120 141 164 166 244 260 278 394 20 42 54 63 70-72 80-81 97 152 164 179
skeletal muscle skeletal muscle skeletal muscle spinal cord adult spleen	Clontech Clontech null Clontech	SKM002 SKMS03 SKMS04 SPC001	322-323 327 341 451 464 246 49 88 6 12 22-23 48 60 70-72 80 93 101-102 104- 106 114 125 135 138-139 143-144 148 162- 163 167 176 196 200 202-204 242-243 264 270 283 288 294 302 321 326 329 333 336 343 345 349 352 354 382 424-425 436 457 2 98 120 141 164 166 244 260 278 394 20 42 54 63 70-72 80-81 97 152 164 179 202 214 238 246 256 311 321 341 353 356
skeletal muscle skeletal muscle skeletal muscle spinal cord adult spleen stomach	Clontech Clontech null Clontech Clontech Clontech	SKM002 SKMS03 SKMS04 SPC001 SPC001	322-323 327 341 451 464 246 49 88 6 12 22-23 48 60 70-72 80 93 101-102 104- 106 114 125 135 138-139 143-144 148 162- 163 167 176 196 200 202-204 242-243 264 270 283 288 294 302 321 326 329 333 336 343 345 349 352 354 382 424-425 436 457 2 98 120 141 164 166 244 260 278 394 20 42 54 63 70-72 80-81 97 152 164 179 202 214 238 246 256 311 321 341 353 356 365 403-404 433
skeletal muscle skeletal muscle skeletal muscle spinal cord adult spleen	Clontech Clontech null Clontech	SKM002 SKMS03 SKMS04 SPC001	322-323 327 341 451 464 246 49 88 6 12 22-23 48 60 70-72 80 93 101-102 104- 106 114 125 135 138-139 143-144 148 162- 163 167 176 196 200 202-204 242-243 264 270 283 288 294 302 321 326 329 333 336 343 345 349 352 354 382 424-425 436 457 2 98 120 141 164 166 244 260 278 394 20 42 54 63 70-72 80-81 97 152 164 179 202 214 238 246 256 311 321 341 353 356 365 403-404 433 1 14 17 23 47 57-58 62 66 70-72 80 101
skeletal muscle skeletal muscle skeletal muscle spinal cord adult spleen stomach	Clontech Clontech null Clontech Clontech Clontech	SKM002 SKMS03 SKMS04 SPC001 SPC001	322-323 327 341 451 464 246 49 88 6 12 22-23 48 60 70-72 80 93 101-102 104- 106 114 125 135 138-139 143-144 148 162- 163 167 176 196 200 202-204 242-243 264 270 283 288 294 302 321 326 329 333 336 343 345 349 352 354 382 424-425 436 457 2 98 120 141 164 166 244 260 278 394 20 42 54 63 70-72 80-81 97 152 164 179 202 214 238 246 256 311 321 341 353 356 365 403-404 433 1 14 17 23 47 57-58 62 66 70-72 80 101 117 134 151 165 187 194 201 220 241 243
skeletal muscle skeletal muscle skeletal muscle spinal cord adult spleen stomach	Clontech Clontech null Clontech Clontech Clontech	SKM002 SKMS03 SKMS04 SPC001 SPC001	322-323 327 341 451 464 246 49 88 6 12 22-23 48 60 70-72 80 93 101-102 104- 106 114 125 135 138-139 143-144 148 162- 163 167 176 196 200 202-204 242-243 264 270 283 288 294 302 321 326 329 333 336 343 345 349 352 354 382 424-425 436 457 2 98 120 141 164 166 244 260 278 394 20 42 54 63 70-72 80-81 97 152 164 179 202 214 238 246 256 311 321 341 353 356 365 403-404 433 1 14 17 23 47 57-58 62 66 70-72 80 101 117 134 151 165 187 194 201 220 241 243 249 278 282 294 337 346 351 353-354 381
skeletal muscle skeletal muscle skeletal muscle spinal cord adult spleen stomach	Clontech null Clontech Clontech Clontech Clontech Clontech	SKM002 SKMS03 SKMS04 SPC001 SPC001 SPLc01 STO001	322-323 327 341 451 464 246 49 88 6 12 22-23 48 60 70-72 80 93 101-102 104- 106 114 125 135 138-139 143-144 148 162- 163 167 176 196 200 202-204 242-243 264 270 283 288 294 302 321 326 329 333 336 343 345 349 352 354 382 424-425 436 457 2 98 120 141 164 166 244 260 278 394 20 42 54 63 70-72 80-81 97 152 164 179 202 214 238 246 256 311 321 341 353 356 365 403-404 433 1 14 17 23 47 57-58 62 66 70-72 80 101 117 134 151 165 187 194 201 220 241 243 249 278 282 294 337 346 351 353-354 381 396 424 430 434
skeletal muscle skeletal muscle skeletal muscle spinal cord adult spleen stomach	Clontech Clontech null Clontech Clontech Clontech	SKM002 SKMS03 SKMS04 SPC001 SPC001	322-323 327 341 451 464 246 49 88 6 12 22-23 48 60 70-72 80 93 101-102 104- 106 114 125 135 138-139 143-144 148 162- 163 167 176 196 200 202-204 242-243 264 270 283 288 294 302 321 326 329 333 336 343 345 349 352 354 382 424-425 436 457 2 98 120 141 164 166 244 260 278 394 20 42 54 63 70-72 80-81 97 152 164 179 202 214 238 246 256 311 321 341 353 356 365 403-404 433 1 14 17 23 47 57-58 62 66 70-72 80 101 117 134 151 165 187 194 201 220 241 243 249 278 282 294 337 346 351 353-354 381

			249 258-259 263-264 284 289 292 298 302 309 311 314 319-320 322-323 329 333 336 341 352 360 371 412 417 440 447 467 472- 473
thymus	Clontech	THMc02	9-10 15 17 24-25 27-28 34 38 40 43 49 57- 58 68 74 77 81 87 94-95 98 104-108 110 115-116 128 136-137 143 146 148-151 158 160 165 197 200 210-211 215 221-222 232 235 241 243 245 248 252 269 278 281 286 288-289 292 302 312 321 325 327 329 331 333 338 345 350 365 378 383 387 412 428 439-440 446 451-452 460 465 469-473
thyroid gland	Clontech	THR001	1 4-5 8-9 11-12 14-15 17 19 21-25 27 34 40 46 49 54-55 57-59 61-62 66-68 70-72 80-81 85 93 97-98 102-108 116 119 121-122 124 126-133 141-142 144 146 150-151 155 162-166 169 171 175-176 178-181 187-190 202-205 208 214-215 218-219 226 232 237-239 244 246-247 250-252 257-258 260 263-264 267 270-271 277 279 282-284 287-288 292 294 297 300 302-304 307-308 310-311 313 317 322-323 325 333 336-337 341 346 356 358 401 405-406 408-409 436 461
trachea .	Clontech	TRC001	17 23 34 90 93 108 142 151 238 240 246 259 266 333 412 472-473
uterus	Clontech	UTR001	18 20 30-31 50 52 114 125 158 164 168 182 198 206 210 248 254-255 260 273 283 304 311 325 365 383 421 423

The 16 tissue/mRNAs and their vendor sources are as follows: 1) Normal adult brain mRNA (Invitrogen), 2) Normal adult kidney mRNA (Invitrogen), 3) Normal fetal brain mRNA (Invitrogen), 4) Normal adult liver mRNA (Invitrogen), 5) Normal fetal kidney mRNA (Invitrogen), 6) Normal fetal liver mRNA (Invitrogen), 7) normal fetal skin mRNA (Invitrogen), 8) human adrenal gland mRNA (Clontech), 9) Human bone marrow mRNA (Clontech), 10) Human leukemia lymphoblastic mRNA (Clontech), 11) Human thymus mRNA (Clontech), 12) human lymph node mRNA (Clontech), 13) human so\spinal cord mRNA (Clontech), 14) human thyroid mRNA (Clontech), 15) human esophagus mRNA (BioChain), 16) human conceptional umbilical cord mRNA (BioChain).

TABLE 2

NO: No. 445 gi4151328 445 gi4151330	Homo sapiens Homo sapiens	high-risk human papilloma viruses E6 oncoproteins targeted protein E6TP1	2344	Identity
445 gi4151330		oncorroteins targeted protein F6TP!		48
445 gi4151330	Va-a engine	1 checking Benea biorent Folia		
gi4151330	U coniens	alpha mRNA, complete cds.		1
	nomo sapiens	high-risk human papilloma viruses E6	1694	59
		oncoproteins targeted protein E6TP1 beta		
		mRNA, complete cds.		
445 gi2555183	Rattus	SPA-1 like protein p1294	2324	48
0	norvegicus	·		
446 gi13517972	Homo sapiens	PR-domain containing protein 17 mRNA,	2496	100
١		complete cds.		
446 gi10434545	Homo sapiens	cDNA FLJ12827 fis, clone	2496	100
"		NT2RP2002939, weakly similar to ZINC		
		FINGER PROTEIN 136.		
446 gi13623607	Homo sapiens	, zinc finger protein 136 (clone pHZ-20),	710	42
3.4		clone MGC:12711, mRNA, complete cds.	1	
447 gi6093239	Homo sapiens	mRNA; cDNA DKFZp434O0515 (from	1054	100
		clone DKFZp434O0515).		
447 gi3522970	Homo sapiens	Trio mRNA, complete cds.	216	23
447 AAW27227		Human TRIO phosphoprotein.	216	23
448 gi7022890	Homo sapiens	cDNA FLJ10700 fis, clone	2838	96
5022050	110,110 0=p10,110	NT2RP3000665.		1
448 gi10438668	Homo sapiens	cDNA: FLJ22327 fis, clone HRC05572.	1333	100
448 gi7020045	Homo sapiens	cDNA FLJ20140 fis, clone COL07182.	1074	79
449 gi6102903	Homo sapiens	mRNA; cDNA DKFZp566D244 (from	2601	99
610102703	Trouse supress	clone DKFZp566D244); partial cds.	1	1
449 gi10434000	Homo sapiens	cDNA FLJ12485 fis, clone	1907	100
61,043,000	Tromo supiens	NT2RM2000420.	1	1.00
449 gi10437387	Homo sapiens	cDNA: FLJ21308 fis, clone COL02131.	1519	69
450 gi7670836	Homo sapiens	hepatocellular carcinoma-associated	3101	99
817070030	Tomo supiens	antigen 66 (HCA66) mRNA, complete	3.01	"
		cds.		
450 gi7959764	Homo sapiens	PRO1289	935	100
450 gj927708	Saccharomyce	Ydr449cp; CAI: 0.18	288	32
3527.100	s cerevisiae			
451 gi7020902	Homo sapiens	cDNA FLJ20657 fis, clone KAT01069.	3231	99
451 gil1037252		NPL4	3156	96
6111037232	norvegicus	11121	3.50] "
451 gi10434779		cDNA FLJ12984 fis, clone	2812	99
6.1045 777	Tromo suprems	NT2RP3000047, weakly similar to NPL4		"
		PROTEIN.	<u> </u>	
452 gi13160469	Homo sapiens	WDR13 protein (WDR13) gene,	1063	94
432 . Billio40	Tiomo sapiens	complete cds.	1003	[7
452 gi12044400	Homo sapiens	WDR13 protein (WDR13) mRNA,	1063	94 .
51120777700	Tronto suprems	complete cds.	1005	'
452 gi13751862	Mus musculus	WD-repeat protein	1058	93
453 gi12619286		mRNA for spinal cord-derived protein	1133	100
g(12017260	Tromo sapiens	FI58G, complete cds.	1133	100
453 gi7638241	Homo sapiens	mesenchymal stem cell protein DSC92	1122	100
453 gi7638241	1101110 Sapieits	mRNA, complete cds.	1133	100
452 -:12004540	Vome coniera	, mesenchymal stem cell protein DSC92,	1122	100
453 gi12804543	Homo sapiens		1133	100
464 :1006000	17	clone MGC:2824, mRNA, complete cds.	2000	100
454 gi13279287	Homo sapiens	, clone IMAGE:3633354, mRNA, partial	2066	100
100	+	cds.		
454 gi5052586	Drosophila	BcDNA.GH08385	334	25
	melanogaster		122	
454 gi10433073	Homo sapiens	cDNA FLJ11749 fis, clone	190	26

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
			HEMBA 1005558, weakly similar to NUCLEAR PROTEIN SNF7.		
455	gi7019840	Homo sapiens	cDNA FLJ20018 fis, clone ADSE00909.	1698	99
455	gi13938166	Homo sapiens	, clone MGC:12617, mRNA, complete cds.	1630	98
455	gi9280376	Homo sapiens	ancient conserved domain protein 3 (ACDP3) mRNA, complete cds.	1271	90
456	gi7020190	Homo sapiens	cDNA FLJ20232 fis, clone COLF5593.	1487	100
456	gi14249896	Homo sapiens	, clone MGC:15774, mRNA, complete cds.	1479	99
456	gi9188416	Homo sapiens	Novel human gene mapping to chomosome 22.	1479	99
457	AAW75093	Homo sapiens	Human secreted protein encoded by gene 37 clone HFVGS85.	369	100
457 .	gi8895089	Homo sapiens	protein x 013 mRNA, complete cds.	145	41
457	gi14250569	Homo sapiens	, protein x 013, clone MGC:3073, mRNA, complete cds.	145	41
458	gi7020228	Homo sapiens	cDNA FLJ20257 fis, clone COLF7231.	1169	100
458	gi7528184	Drosophila melanogaster	bicoid-interacting protein BIN3	389	45
459	gi11345384	Homo sapiens	vacuolar protein sorting protein 18 (VPS18) mRNA, complete cds.	5102	100
459	AAW48303	Homo sapiens	Amino acid sequence of human deep orange protein.	2555	100
459	gi2832850	Drosophila melanogaster	EG:171E4.1	1316	35
460	gi6966967	Homo sapiens	mRNA for dipeptidyl-peptidase III (DPP3 gene).	3814	99
460	gi13938201	Homo sapiens	, dipeptidylpeptidase III, clone MGC:15061, mRNA, complete cds.	3811	99
460	AAB67571	Homo sapiens	Amino acid sequence of a human hydrolytic enzyme HYENZ3.	3807	99
461	AAY53020	Homo sapiens	Human secreted protein clone qb56_19 protein sequence SEQ ID NO:46.	657	100
461	AAY59788	Homo sapiens	Human normal ovarian tissue derived protein 65.	618	100
461	AAG04028	Homo sapiens	Human secreted protein, SEQ ID NO: 8109.	442	72
462	gi13021843	Homo sapiens	polyadenylate binding protein-interacting protein 2 mRNA, complete cds.	679	100
462	gi12052806	Homo sapiens	mRNA; cDNA DKFZp564F163 (from clone DKFZp564F163); complete cds.	675	99
462	gi7106826	Homo sapiens	HSPC218	673	99
463	gi7023258	Homo sapiens	cDNA FLJ10914 fis, clone OVARC1000212.	1067	100
464	gi7023258	Homo sapiens	cDNA FLJ10914 fis, clone OVARC1000212.	649	72
465	gi7022147	Homo sapiens	cDNA FLJ10233 fis, clone HEMBB1000266.	3464	100
465	gi12224837	Homo sapiens	mRNA; cDNA DKFZp547K202 (from clone DKFZp547K202).	3464	100
465	AAY99662	Homo sapiens	Human GTPase associated protein-13.	3464	100
466	gi7582304	Homo sapiens	BM-016	584	100
466	AAW85610	Homo sapiens	Secreted protein clone eh80_1.	330	97
466	AAW78199	Homo sapiens	Human secreted protein encoded by gene 74 clone HGBAC11.	330	97
467	gi7018410	Homo sapiens	mRNA; cDNA DKFZp566K023 (from clone DKFZp566K023).	1010	100
467	gi9049987	Rattus	X2CR1 protein	268	81

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
		norvegicus			
468	gi8317213	Homo sapiens	histone acetyltransferase (MOF) mRNA, partial cds.	1625	100
468	gi10433157	Homo sapiens	cDNA FLJ11810 fis, clone HEMBA1006347, moderately similar to MALES-ABSENT ON THE FIRST	1625	100
468	gi10436400	Homo sapiens	PROTEIN (EC 2.3.1). cDNA FLJ14040 fis, clone HEMBA1005513, weakly similar to MALES-ABSENT ON THE FIRST PROTEIN (EC 2.3.1).	1613	99
469	AAY76072	Homo sapiens	Human skin cell protein, SEQ ID NO:327.	668	100
469	AAB56011	Homo sapiens	Skin cell protein, SEQ ID NO: 327.	668	100
470	gi29481	Homo sapiens	Human erythrocyte 2,3- bisphosphoglycerate mutase mRNA EC 2.7.5.4.	1362	100
470	gi 179527	Homo sapiens	Human 2,3-bisphosophoglycerate mutase (BPGM) gene, exon 3.	1362	100
470	AAB11959	Homo sapiens	Glycated human erythrocyte bisphosphoglycerate mutase (BPGM).	.1362	100
471	gi6841472	Homo sapiens	HSPC125	892 .	100
471	gi12001966	Homo sapiens	clone 015g09 My013 protein mRNA, complete cds.	892	100
471	gi9624483	Homo sapiens	HRPAP20 short form mRNA, complete cds.	640	72
472	gi9367763	Homo sapiens	mRNA for zinc finger protein Cezanne (CEZANNE gene).	2580	100
472	gi6102920	Homo sapiens	mRNA; cDNA DKFZp434H0717 (from clone DKFZp434H0717); partial cds.	2197	100
472	gi7332054	Caenorhabditis elegans	contains similarity to tumor necrosis factors	126	25
473	gi8489813	Homo sapiens	DJ963K23.2 mRNA, complete cds.	1255	100
473	AAB43861	Homo sapiens	Human cancer associated protein sequence SEQ ID NO:1306.	1255	100
473	gi9858803	Mus musculus	Zfp228	1090	91 ·
474	gi7020223	Homo sapiens	cDNA FLJ20254 fis, clone COLF6926.	2278	100
474	AAY25743	Homo sapiens	Human secreted protein encoded from gene 33.	917	100
474	AAY76166	Homo sapiens	Human secreted protein encoded by gene 43.	724	94
475	gi14042066	Homo sapiens	cDNA FLJ14503 fis, clone NT2RM1000252, weakly similar to H.sapiens E-MAP-115 mRNA.	159	26
475	gi7270600	Arabidopsis thaliana	trichohyalin like protein	156	25
475	gi180195	Homo sapiens	Human aorta caldesmon mRNA, complete cds.	145	25
476	gi11066250	Homo sapiens	presenilins associated rhomboid-like protein (PARL) mRNA, complete cds.	2030	100
476	gi13177766	Homo sapiens	, Similar to presenilins associated rhomboid-like protein, clone MGC:4756, mRNA, complete cds.	1107	99
476	gi7959883	Homo sapiens	PRO2207	986	100
477	AAY91941	Homo sapiens	Human chaperone protein 2 (HCHP-2).	1977	100
477	gi7019854	Homo sapiens	cDNA FLJ20027 fis, clone ADSE01901.	1965	99
477	gi6567172	Mus musculus	mDj10	1863	93
478	gi13937971	Homo sapiens	; Similar to RIKEN cDNA 1110005A23 gene, clone MGC:14726, mRNA,	1040	100

SEQ ID NO:	Accession No.	Species	Description .	Score	% Identity
110.	110.		complete cds.		
478	gi13940310	Homo sapiens	HCC-1 gene.	1040	100
478	AAB36609	Homo sapiens	Human FLEXHT-31 protein sequence SEQ ID NO:31.	1040	100
479	gi11065999	Homo sapiens	neuronal calcium binding protein NECAB3 mRNA, complete cds.	1889	99
479	gi10798741	Homo sapiens	XB51 mRNA for X11L-binding protein 51, complete cds.	654	99
479	gi10798743	Mus musculus	X11L binding protein 51	1079	86
480	gi6094684	Homo sapiens	PAC clone RP1-278D1 from X, complete sequence.	3056	92
480	gi10435614	Homo sapiens	cDNA FLJ13568 fis, clone PLACE1008368, weakly similar to RING CANAL PROTEIN.	1847	100
480	gi7023516	Homo sapiens	cDNA FLJ11078 fis, clone PLACE1005102, weakly similar to RING CANAL PROTEIN.	1208	42
481	gi7020424	Homo sapiens	cDNA FLJ20369 fis, clone HEP19364.	2727	100
481	gi1110599	Mus sp.	semaphorin homolog=M-Sema F	2653	86
481	AAB88485	Homo sapiens	Human membrane or secretory protein clone PSEC0078.	1774	100
482	gi4679028	Homo sapiens	HSPC021	1930	100
482	gi5106781	Homo sapiens	HSPC025	1930	100
482	gi12654535	Homo sapiens	, HSPC025, clone MGC:735, mRNA, complete cds.	1930	100
483	gi1145789	Rattus norvegicus	neuroligin 2	4417	98
483	gi7960135	Homo sapiens	neuroligin 3 isoform gene, complete cds, alternatively spliced.	2736	65
483	gi7960131	Homo sapiens	neuroligin 3 isoform HNL3 mRNA, complete cds, alternatively spliced.	2729	65
484	gi14250554	Homo sapiens	, hexokinase 1, clone MGC:1724, mRNA, complete cds.	4725	99
484	gi2873349	Homo sapiens	hexokinase I (HK1) gene, exon 18, complete cds, alternatively spliced.	4725	99
484	gi184021	Homo sapiens	Human hexokinase 1 (HK1) mRNA, complete cds.	4718	99
485	gi8453103	Homo sapiens	zinc finger protein mRNA, complete cds.	3726	100
485	gi13752754	Homo sapiens	zinc finger 1111 mRNA, complete cds.	1689	56
485	gi10436789	Homo sapiens	cDNA FLJ14345 fis, clone THYRO1001189, weakly similar to ZINC FINGER PROTEIN 91.	1683	56
486	AAB56937	Homo sapiens	Human prostate cancer antigen protein sequence SEQ ID NO:1515.	2341	100
486	gi12804453	Homo sapiens	, Similar to Tu translation elongation factor, mitochondrial, clone MGC:1592, mRNA, complete cds.	2326	100
486	gi899285	Homo sapiens	H.sapiens mRNA for elongations factor Tu-mitochondrial.	2326	100
487	gi9910111	Homo sapiens	myosin X (MYO10) mRNA, complete cds.	10727	99
487	gi6996558	Mus musculus		10089	93
487	gi7108753	Homo sapiens	myosin X (MYO10) mRNA, partial cds.	8029	99
488	gi7688687	Homo sapiens		1935	100
488	gi14042251	Homo sapiens	cDNA FLJ14611 fis, clone NT2RP1000988.	1935	100
488	AAY66671	Homo sapiens	Membrane-bound protein PRO1134.	1935	100
489	gi202215	Mus musculus		2387	100
489	gi14328047	Homo sapiens	, tubulin alpha 4, clone MGC:2379,	2387	100

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
			mRNA, complete cds.		
489	gi1333692	Macaca fascicularis	alpha-tubulin (ATG-initiation codon missing)	2382	100
490	gi5912034	Homo sapiens	mRNA; cDNA DKFZp434N0535 (from clone DKFZp434N0535); partial cds.	6810	99
490	gi5912239	Homo sapiens	mRNA; cDNA DKFZp434O225 (from clone DKFZp434O225); partial cds.	3442	99
490	gi3292939	Drosophila melanogaster	Additional sex combs	295	39
491	gi5912034	Homo sapiens	mRNA; cDNA DKFZp434N0535 (from clone DKFZp434N0535); partial cds.	5941	99
491	gi5912239	Homo sapiens	mRNA; cDNA DKFZp434O225 (from clone DKFZp434O225); partial cds.	2573	99
491	gi3292939	Drosophila melanogaster	Additional sex combs	295	39
492	AAY68778	Homo sapiens	Amino acid sequence of a human phosphorylation effector PHSP-10.	2463	99
492	gi479173	Homo sapiens	H.sapiens nek3 mRNA for protein kinase.	2417	99
492	gi13529320	Mus musculus	Similar to NIMA (never in mitosis gene a)-related expressed kinase 3	1887	73
493	gi13539686	Homo sapiens	protein kinase C and casein kinase substrate 1 (PACSIN1) mRNA, complete cds.	2365	100
493	gi728604	Mus musculus	PACSIN	2250	95
493	gi4324452	Rattus norvegicus	syndapin l	2250	95
494	gi7023749	Homo sapiens	cDNA FLJ11220 fis, clone PLACE1008129.	3994	. 100
494	gi10433501	Homo sapiens	cDNA FLJ12104 fis, clone HEMBB1002697.	2829	100
494	gi5788108	Homo sapiens	PAC clone RP5-1087M19 from 7q11.23- q21.1, complete sequence.	757	63
495	AAB54375	Homo sapiens	Human pancreatic cancer antigen protein sequence SEQ ID NO:827.	2897	99
495	AAY57923	Homo sapiens	Human transmembrane protein HTMPN-47.	2724	98
495	AAW88628	Homo sapiens	Secreted protein encoded by gene 95 clone HPWAN23.	2686	98
496	gi7959788	Homo sapiens	PRO1635	317	100
496	AAW74852	Homo sapiens	Human secreted protein encoded by gene 124 clone HPCAD23.	143	100
497	gi7707424	Homo sapiens	mRNA for syntaxin 18, complete cds.	1705 1488	83
498	gi1613858	Homo sapiens	Human zinc finger protein zfp47 (zf47) mRNA, partial cds.		
498	gi13938633	Mus musculus	RIKEN cDNA 2810435N07 gene	1318	58
498	gi9837564	Mus musculus	SCAN-KRAB-zinc finger protein	1242	
499	AAY27795	Homo sapiens	Human secreted protein encoded by gene No. 79.	1539	99
499	gi10436317	Homo sapiens	cDNA FLJ13986 fis, clone Y79AA1001923, weakly similar to Homo sapiens F-box protein Fbx22 (FBX22) gene.	1370	100
499	gi6164747	Homo sapiens	F-box protein Fbx22 (FBX22) gene, partial cds.	391	93
500	gi3150052	Homo sapiens	mRNA, complete cds.	4455	100
500	gi14280050	Homo sapiens	cds.	382	24
500	gi12718237	Neurospora	related to TGF beta receptor associated	174	31

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SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
		crassa ·	protein 1		
501	gi7023051	Homo sapiens	cDNA FLJ10796 fis, clone NT2RP4000648, weakly similar to TRANS-ACTING TRANSCRIPTIONAL PROTEIN ICPO.	3360	99
501	gi9651170	Homo sapiens	cell cycle checkpoint protein CHFR mRNA, complete cds.	2491	96
501	AAB20219	Homo sapiens	Human Chfr (checkpoint with FHA and ring finger) protein.	2491	96
502	gi7329074	Homo sapiens	collagen type V alpha 3 chain (COL5A3) mRNA, complete cds.	9671	100
502	gi8568094	Rattus norvegicus	alpha 4 type V collagen	8038	82
502	gi7329072	Mus musculus	collagen type V alpha 3 chain	7970	82
503	gi12654687	Homo sapiens	, clone MGC:2616, mRNA, complete cds.	1161	100
503	gi7769617	Mus musculus	TCE2	1050	89
504	gi12654687	Homo sapiens	, clone MGC:2616, mRNA, complete cds.	1140	96
504	gi7769617	Mus musculus	TCE2	1029	86 .
505	gi12654687	Homo sapiens	, clone MGC:2616, mRNA, complete cds.	654	100
505	gi7769617	Mus musculus	TCE2	629	92
506	gi14249942	Homo sapiens	, Similar to RIKEN cDNA 0610008P16 gene, clone MGC:15937, mRNA, complete cds.	1609	100
506	AAB56487	Homo sapiens	Human prostate cancer antigen protein sequence SEQ ID NO:1065.	1167	98
506	gi2828262	Bos taurus	aralkyl acyl-CoA:amino acid N- acyltransferase	597	40
507	gi7688987	Homo sapiens	uncharacterized bone marrow protein BM046	1295	100
507	AAB64387	Home sapiens	Amino acid sequence of human intracellular signalling molecule INTRA19.	1202	94
507	gi9437511	Homo sapiens	BM024	1045	98
508	AAB18979	Homo sapiens	Amino acid sequence of a human transmembrane protein.	1203	100
508	gi6808196	Homo sapiens	mRNA; cDNA DKFZp434P1018 (from clone DKFZp434P1018); partial cds.	938	100
508	gi13960126	Homo sapiens	, Similar to leucine-rich neuronal protein, clone MGC:4126, mRNA, complete cds.	845	100
509	gi13938527	Homo sapiens	, Similar to RIKEN cDNA 2810002N01 gene, clone MGC:2562, mRNA, complete cds.	1048	100
509	AAY35994	Homo sapiens	Extended human secreted protein sequence, SEQ ID NO. 379.	1032	98
509	AAG00345	Homo sapiens	Human secreted protein, SEQ ID NO: 4426.	619	98
510	gi773387	Neurospora crassa	Restriction enzyme inactivation of met-10 complementation in this region. Sequence similarity to S. cerevisiae chromosome VIII cosmid 9205, accession no. U10556 CDS residues 22627-24126	536	35
510	gi487945	Saccharomyce s cerevisiae	Yhr070wp	528	49
510	AAG02508	Homo sapiens	Human secreted protein, SEQ ID NO: 6589.	324	100
511	gi11493195	Homo sapiens	mRNA for LB1 protein.	2614	99
511	gi10434688	Homo sapiens	cDNA FLJ12920 fis, clone NT2RP2004594.	2604	99
511	gi12053201	Homo sapiens	mRNA; cDNA DKFZp434A1031 (from	2604	99

SEQ ID NO:	Accession No.	. Species	Description	Score	% Identity
110.	140.		clone DKFZp434A1031); complete cds.		lucinity
512	AAW75106	Homo sapiens	Human secreted protein encoded by gene 50 clone HHSDZ57.	471	100 .
512	AAY59689	Homo sapiens	Secreted protein 26-44-1-B5-CL3_1.	471	100
512	AAY48331	Homo sapiens	Human prostate cancer-associated protein 28.	471	100
514	AAW67888	Homo sapiens	Human secreted protein encoded by gene 82 clone HSKHL65.	921	92
514	gi13436110	Homo sapiens	, Similar to RIKEN cDNA 2310034L04 gene, clone MGC:11061, mRNA, complete cds.	150	28
514	AAY53052	Homo sapiens	Human secreted protein clone df202_3 protein sequence SEQ ID NO:110.	132	33
515	gi7020259	Homo sapiens	cDNA FLJ20276 fis, clone HEP02437.	5378	100
515	gi10432807	Homo sapiens	cDNA FLJ11534 fis, clone HEMBA1002679.	3024	99
515	gi9916	Plasmodium falciparum	liver stage antigen	399	23
516	AAB67448	Homo sapiens	Amino acid sequence of a human chaperone polypeptide.	1190	99 .
516	gi13477189	Homo sapiens	, Similar to RIKEN cDNA 1300007M11 gene, clone MGC:12943, mRNA, complete cds.	1182	99
516	AAG03527	Homo sapiens	Human secreted protein, SEQ ID NO: 7608.	389	98
517	gi7023782	Homo sapiens	cDNA FLJ11240 fis, clone PLACE1008568.	2796	100
517	AAB08869	Homo sapiens	Amino acid sequence of a human secretory protein.	2792	99
517.	AAB23626	Homo sapiens	Human secreted protein SEQ ID NO: 52.	2792	99
518	gi6460009	Deinococcus radiodurans	citrate lyase, beta subunit	211	30
518	gi14025765	Mesorhizobiu m loti	citrate lyase beta-subunit	324	31
518	gi14024477	Mesorhizobiu m loti	Citrate lyase beta chain (acyl lyase subunit); CitE	316	33
519	gi14041831	Homo sapiens	cDNA FLJ14357 fis, clone HEMBA1000005, highly similar to DNAJ PROTEIN HOMOLOG MTJ1.	2873	100
5,19	AAB67447	Homo sapiens	Amino acid sequence of a human chaperone polypeptide.	2481	99
519	gi473847	Mus musculus	dnaJ-like protein	2413	84
520	gi7669968	Homo sapiens	mRNA; cDNA DKFZp761G0313 (from clone DKFZp761G0313).	789	100
520	gi4586315	Homo sapiens	ORCTL3 mRNA for organic-cation transporter like 3, complete cds.	348	38
520	gi4835384	Homo sapiens	DNA, DLEC1 to ORCTL4 gene region, section 1/2 (DLEC1, ORCTL3, ORCTL4 genes, complete cds).	348	38
521	gi7959805	Homo sapiens	PRO0823	344	100
522	gi10434341	Homo sapiens	cDNA FLJ12691 fis, clone NT2RM4002571, weakly similar to H.sapiens mRNA for UDP- GaINAc:polypeptide N- acetylgalactosaminyltransferase (T2).	2605	89
522	gi10436305	Homo sapiens	cDNA FLJ13977 fis, clone Y79AA1001603, weakly similar to POLYPEPTIDE N- ACETYLGALACTOSAMINYLTRANS FERASE (EC 2.4.1.41).	1631	99

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
522	gi971461	Homo sapiens	H.sapiens mRNA for UDP- GalNAc:polypeptide N- acetylgalactosaminyltransferase (T2).	1386	50
523	gi11493500	Homo sapiens	PRO2979	477	100
523	gi38163	Pan troglodytes	A-gamma-globin	477	100
523	gi176779	Pan troglodytes	gamma-2 globin	477	100
524	gi5262582	Homo sapiens	mRNA; cDNA DKFZp434K063 (from clone DKFZp434K063); partial cds.	3782	99
524	gi10438230	Homo sapiens	cDNA: FLJ21993 fis, clone HEP06576.	1416	100
524	AAY21842	Homo sapiens	Human signal peptide-contianing protein (SIGP) (clone ID 1273453).	1416	100
525	gi1928886	Rattus norvegicus	lin-10 protein homolog	2199	97
525	gi10433467	Homo sapiens	cDNA FLJ12076 fis, clone HEMBB1002442, weakly similar to LIN- 10 PROTEIN.	483	98
525	gi5824587	Caenorhabditis elegans	T01G9.2b	668	37
526	gi1679607	Mus musculus	myosin-I	4206	84
526	gi1924940	Homo sapiens	H.sapiens mRNA for myosin-IE.	4115	99
526	gi65324	Gallus gallus	brush border myosin IB	3812	76
527	AAB63419	Homo sapiens	Human breast cancer associated antigen protein sequence SEQ ID NO:781.	641	99
528	gi13649967	Homo sapiens	fovea-associated SH3 domain binding protein (FASH3) mRNA, complete cds.	558	.100
528	gi13539561	Homo sapiens	mRNA for SH3BGRL2 protein.	558	100
528	gi5042302	Mus musculus	sh3bgr protein	365	64
529	gi10436540	Homo sapiens	cDNA FLJ14154 fis, clone NT2RM1000341.	1151	99
529	gi13436011	Mus musculus	RIKEN cDNA 1200013P24 gene	1139	97
529	gi1592161	Methanococcu s jannaschii	ribosomal protein S18 alanine acetyltransferase	109	36
530	gi3135314	Homo sapiens	chromosome 7q22 sequence, complete sequence.	911	100
530	gi6752287	Homo sapiens	Novel human gene mapping to chomosome X.	281	51
531	gi14042818	Homo sapiens	cDNA FLJ14937 fis, clone PLACE1010231, weakly similar to CELL SURFACE GLYCOPROTEIN EMR1 PRECURSOR.	2548	97
531	gi2117161	Homo sapiens	H.sapiens mRNA for HE6 Tm7 receptor.	1366	52
531	AAW36903	Homo sapiens	Human epididymis-specific receptor protein.	1366	52
532	gi7417372	Homo sapiens	intracellular hyaluronan-binding protein mRNA, complete cds.	2175	99
532	gi7110497	Mus musculus	intracellular hyaluronan-binding protein p57	1862	85
532	gi3403154	Homo sapiens	Human Ki-1/57 intracellular antigen mRNA, partial cds.	1591	98
533	gi10436645	Homo sapiens	cDNA FLJ14235 fis, clone NT2RP4000167.	1585	82
533	gi7020976	Homo sapiens	cDNA FLJ20707 fis, clone KAIA1223.	2195	84
533	gi13276619	Homo sapiens	mRNA; cDNA DKFZp76110112 (from clone DKFZp76110112).	1444	99
534	gi438880	Rattus norvegicus	tropomyosin	1186	99
534	gi2978558	Xenopus	alpha-tropomyosin	1089	89

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
		laevis			
534	gi438882	Rattus norvegicus	tropomyosin	1086	92
535	gi438880	Rattus norvegicus	tropomyosin	1120	93
535	gi9508585	Homo sapiens	tropomyosin isoform mRNA, complete cds.	1105	93
535	gi12653955	Homo sapiens	, Similar to tropomyosin 4, clone MGC:3261, mRNA, complete cds.	1094	91
536	gi6808111	Homo sapiens	mRNA; cDNA DKFZp434O1230 (from clone DKFZp434O1230); partial cds.	439	100
537	gi6807806	Homo sapiens	mRNA; cDNA DKFZp434K031 (from clone DKFZp434K031); partial cds.	3007	100
537	gi13623334	Homo sapiens	, Similar to DKFZP727C091 protein, clone MGC:10677, mRNA, complete cds.	2392 i	100
537	AAY25821	Homo sapiens	Human secreted protein fragment encoded from gene 41.	1967	99 .
538	AAB88413	Homo sapiens	Human membrane or secretory protein clone PSEC0170.	1818	99
538	gi6457342	Homo sapiens	E2IG4 (E2IG4) mRNA, complete cds.	1813	99
538	AAB24026	Homo sapiens	Human PRO1788 protein sequence SEQ IDNO:18.	1813	99
539	gi6572289	Homo sapiens	mRNA for mitochondrial tryptophanyl- tRNA synthetase (WARS2 gene).	1820	100
539	gi13421159	Caulobacter crescentus	tryptophanyl-tRNA synthetase	727	46
539	gil 1992026	Zymomonas mobilis	tryptophanyl-tRNA synthase	721	43
540	gi7106630	Homo sapiens	Novel human mRNA from chromosome 1, clone Z98884, has homology to PERIOD CIRCADIAN PROTEIN 3.	6301	99
540	gi13160925	Homo sapiens	mRNA for period (Drosophila) homolog 3 hPER3, complete cds.	6274	99
540	AAB23266	Homo sapiens	Human circadian rhythm protein Per3 (hPer3).	6274	99
541	gi9621744	Homo sapiens	ferritin heavy chain subunit mRNA, complete cds.	968	100
541	gi12654093	Homo sapiens	, ferritin, heavy polypeptide 1, clone MGC:5580, mRNA, complete cds.	968	100
541	gi12655095	Homo sapiens	, ferritin, heavy polypeptide 1, clone MGC:1749, mRNA, complete cds.	968	100
542	gi4902699	Homo sapiens	Novel human gene mapping to chomosome 13.	2372	57
542	gi2341020	Homo sapiens	PAC clone 248O15 from 13q12-q13, complete sequence.	1447	58
542	gi11907986	Drosophila melanogaster	fry	1054	38
543	gi7582278	Homo sapiens	BM-003	1386	100
543	gi7688983	Homo sapiens	uncharacterized bone marrow protein BM044	1386	100
543	gi1752736	Saccharomyce s cerevisiae	gene required for phosphoylation of oligosaccharides/ has high homology with YJR061w	150	35
544	gi1628401	Homo sapiens	H.sapiens mRNA for leucine-rich primary response protein 1.	3936	98
544	gi940821	Rattus norvegicus	LRPRI	2914	73
544	gi2196560	Schizosacchar omyces pombe	Mis6	223	31

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
545	gi7022824	Homo sapiens	cDNA FLJ10656 fis, clone NT2RP2006038.	1574	99
545	gi6841138	Homo sapiens	HSPC099 mRNA, partial cds.	248	36
545	AAG02788	Homo sapiens	Human secreted protein, SEQ ID NO: 6869.	234	85
546	AAB71914	Homo sapiens	Human ISOM-6.	1142	98
546	gi3876969	Caenorhabditis elegans	Similarity to Brugia peptidylprolyl isomerase (TR:G984562), contains similarity to Pfam domain: PF00076 (RNA recognition motif. (a.k.a. RRM, RBD, or RNP domain)), Score=62.0, E-value=4.2e-15, N=1; PF00160 (Cyclophilin type peptidyl-prolyl cis-trans isomerase), Score=78.1, E-value=3.7e-22, N=1	658	52
546	AAG02246	Homo sapiens	Human secreted protein, SEQ ID NO: 6327.	573	100
547	gi603635	Saccharomyce s cerevisiae	Yel044wp	133	25
548	gi5262665	Homo sapiens	mRNA; cDNA DKFZp564B0769 (from clone DKFZp564B0769); partial cds.	1455	99
548	gi6841172	Homo sapiens	HSPC261	716	99
548	gi12803875	Homo sapiens	, Similar to splicing factor, arginine/serine-rich 4, clone MGC:3920, mRNA, complete cds.	352	33
549	gi7582298	Homo sapiens	BM-013	704	100
549	gi9558483	Ciona savignyi	PEM-3	434	55
549	gi1644450	Caenorhabditis elegans	MEX-3	362	65
550	gi4883433	Homo sapiens	mRNA for membrane transport protein (XK gene).	2148	100
550	gi6502963	Mus musculus	KX antigen	1797	18
550	gi2580580	Homo sapiens	testis-specific XK Related Y (XKRY) mRNA, complete cds.	157	31
551	gi7670746	Homo sapiens	UDP-glucose:glycoprotein glucosyltransferase 1 precursor, mRNA, complete cds.	8075	99
551	gi13275621	synthetic construct	Rat RUGT	7371	91
551	gi7677176	Rattus norvegicus	UDP-glucose glycoprotein:glucosyltransferase precursor	7371	91
552	gi7688985	Homo sapiens	uncharacterized bone marrow protein BM045	390	72
553	gi12655091	Homo sapiens	, AD-003 protein, clone MGC:783, mRNA, complete cds.	1177	100
553	gi6523799	Homo sapiens	adrenal gland protein AD-003 mRNA, complete cds.	1168	99
553	gi7105659	Caenorhabditis elegans	contains similarity to Streptomyces peucetius carminomycin 4-O- methyltransferase (GB:L13453)	425	39
554	gi7582282	Homo sapiens	BM-005	3445	99
554	gi7022933	Homo sapiens	cDNA FLJ10725 fis, clone NT2RP3001214.	3312	100
554	gi10435575	Homo sapiens	cDNA FLJ13534 fis, clone PLACE1006445.	1648	100
555	gi12751374	Homo sapiens	paraoxanase-3 mRNA, partial cds.	1819	99 . '
555	gi1333634	Homo sapiens	paraoxonase 3 (PON3) mRNA, 3' end of cds.	1741	98
555	gi12743899	Oryctolagus	paraoxonase 3	1542	82

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
		cuniculus			
556	gi7022174	Homo sapiens	cDNA FLJ10252 fis, clone HEMBB1000807.	2826	100
556	gi11596985	Homo sapiens	chromosome 14 clone RP11-361H10 map 14q24.3, complete sequence.	559	36
556	gi7020945	Homo sapiens	cDNA FLJ20689 fis, clone KAIA2890.	510	39
557	gi10434683	Homo sapiens	cDNA FLJ12917 fis, clone	2879	99
•			NT2RP2004568, weakly similar to PUTATIVE ATP-DEPENDENT RNA HELICASE C30D11.03.		
557	gi13384106	Homo sapiens	RNA helicase-like protein (RHLP) mRNA, complete cds.	2817	99
557	gi7020811	Homo sapiens	cDNA FLJ20596 fis, clone KAT08049.	2020	99
558	gi4760710	Brassica rapa	SLL2-S9-protein	284	43
558	gi1669601	Arabidopsis thaliana	AR401	280	44
558	gi557805	Saccharomyce s cerevisiae	orf, len: 257, CAI: 0.13	327	34
559	gi13548677	Homo sapiens	MKP-7 mRNA for MAPK phosphatase-7, complete cds.	3418	100
559	gi13990989	Mus musculus	MAP kinase phosphatase-7	3093	90
559	AAB20325	Homo sapiens	Human protein phosphatase and kinase protein-4.	3021	90
560	gi10433965	Homo sapiens	cDNA FLJ 12464 fis, clone NT2RM1000780.	2196	97
560	gi10434795	Homo sapiens	cDNA FLJ12992 fis, clone NT2RP3000149.	2196	97
560	gi10438048	Homo sapiens	cDNA: FLJ21857 fis, clone HEP02294.	2151	94
561	gi10438048	Homo sapiens	cDNA: FLJ21857 fis, clone HEP02294.	2276	97
561	gi10433965	Homo sapiens	cDNA FLJ 12464 fis, clone NT2RM1000780.	2159	94
561	gi10434795	Homo sapiens	cDNA FLJ12992 fis, clone NT2RP3000149.	2159	94
562	gi10433965	Homo sapiens	cDNA FLJ 12464 fis, clone NT2RM1000780.	2443	99
562	gi10434795	Homo sapiens	cDNA FLJ12992 fis, clone NT2RP3000149.	2443	99
562	gi10438048	Homo sapiens	cDNA: FLJ21857 fis, clone HEP02294.	2398	96
563	gi11137965	Homo sapiens	tRNA isopentenylpyrophosphate transferase precursor RNA, complete cds.	2158	100
563	gi7019915	Homo sapiens	cDNA FLJ20061 fis, clone COL01383.	1719	100
563	gi9803035	Caenorhabditis elegans	contains similarity to Pfam domain PF00096 (zf-C2H2), Score=12.0, E- value=1.1, N=1	407	32
564	gi7023103	Homo sapiens	cDNA FLJ10826 fis, clone NT2RP4001100.	2171	100
564	gi10434339	Homo sapiens	cDNA FLJ12690 fis, clone NT2RM4002567.	2171	100
564	gi10433458	Homo sapiens	cDNA FLJ12068 fis, clone HEMBB1002329.	2166	99
565	gi7019829	Homo sapiens	cDNA FLJ20011 fis, clone ADKA03432.	865	100
565	gi10438448.	Homo sapiens	cDNA: FLJ22168 fis, clone HRC00618.	865	100
565	AAG02581	Homo sapiens	Human secreted protein, SEQ ID NO: 6662.	445	98
566	gi11558482	Homo sapiens	mRNA for B-cell lymphoma/leukaemia 11A extra long form (BCL11A-XL gene).	1543	99
566	gi12150278	Homo sapiens	C2H2-type zinc-finger protein mRNA, complete cds.	1039	99 .
566	gi6652688	Mus musculus	C2H2-type zinc finger protein	1033	98

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
567	gi12053249	Homo sapiens	mRNA; cDNA DKFZp434A155 (from clone DKFZp434A155); complete cds.	994	100
567	AAY73435	Homo sapiens	Human secreted protein clone yd73_1 protein sequence SEQ ID NO:92.	994	100
567	AAB43698	Homo sapiens	Human cancer associated protein sequence SEQ ID NO:1143.	752	95
568	gi12053249	Homo sapiens	mRNA; cDNA DKFZp434A155 (from clone DKFZp434A155); complete cds.	752	95
568	AAY73435	Homo sapiens	Human secreted protein clone yd73_1 protein sequence SEQ ID NO:92.	752	95
568	AAB43698	Homo sapiens	Human cancer associated protein sequence SEQ ID NO:1143.	853	100
569	gi8096260	Homo sapiens	gene for Nop10p, complete cds.	344	100
569	gi8096476	Homo sapiens	mRNA for Nop10p, complete cds.	344	100
569	gi14424489	Homo sapiens	, nucleolar protein family A, member 3 (H/ACA small nucleolar RNPs), clone MGC:19486, mRNA, complete cds.	344	100
570	gi11595476	Homo sapiens	mRNA for RPB11b1beta protein (POLR2J2 gene).	633	100
570	AAB58870	Homo sapiens	Breast and ovarian cancer associated antigen protein sequence SEQ 1D 578.	409	100
570	gi11595474	Homo sapiens	mRNA for RPB11b1alpha protein (POLR2J2 gene).	247	97
571	gi7239381	Homo sapiens	guanine nucleotide exchange factor smgGDS (RAP1GDS1) mRNA, alternatively spliced, complete cds.	2995	99
571	gi13111713	Homo sapiens	, RAP1, GTP-GDP dissociation stimulator 1, clone MGC:2897, mRNA, complete cds.	2994	99
571	gi6942013	Homo sapiens	exchange factor smgGDS mRNA, complete cds, alternatively spliced.	2991	99
572	gi12002978	Homo sapiens	mitosin-associated protein MITAPI (MITAPI) mRNA, complete cds.	1736	100
572	gi12043569	Homo sapiens	Nudel mRNA, complete cds.	1736	100
572	gi13775593	Homo sapiens	endooligopeptidase A mRNA, complete cds.	1720	99
573	gi7022325	Homo sapiens	cDNA FLJ10350 fis, clone NT2RM2001131.	1243	100
573	gi12052730	Homo sapiens	mRNA; cDNA DKFZp761F19121 (from clone DKFZp761F19121).	1243	100
573	gi3417386	Mus musculus	microtubule-associated protein, MAP-115	428	48
574	gi7022502	Homo sapiens	cDNA FLJ10458 fis, clone NT2RP1001457, highly similar to Homo sapiens partial mRNA for beta-transducin family protein.	2555	100
574	gi3687833	Xenopus laevis	notchless	2149	82
574	gi12643028	Oryza sativa	Putative Notchless protein homolog	1110	52
575	AAY51115	Homo sapiens	Human HSEC6 protein.	3767	99
575	gi1163174	Rattus norvegicus	similar to yeast Sec6p, Swiss-Prot Accession Number P32844; similar to mammalian B94, Swiss-Prot Accession Number Q03169; Method: conceptual translation supplied by author	3606	94
575	AAB49655	Homo sapiens	Human SEC7 protein sequence SEQ ID 14.	2737	89
576	gi7020303	Homo sapiens	cDNA FLJ20300 fis, clone HEP06465.	1697	99
576	AAB67575	Homo sapiens	Amino acid sequence of a human	759	47
	<u></u>	1	hydrolytic enzyme HYENZ7.		<u></u>

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
576	gi10434892	Homo sapiens	cDNA FLJ13055 fis, clone NT2RP3001538, weakly similar to HYPOTHETICAL 39.0 KD PROTEIN T28D9.3 IN CHROMOSOME II.	755	47
577	AAR15222	Homo sapiens	Chronic myelogenous leukaemia-derived myeloid-related protein.	513	100
577	gi32402	Homo sapiens	Human mRNA for HP-1, a member of the corticostatin/defensin family.	493	100
577	gi181527	Homo sapiens	Human neutrophil peptide (defensin) 1 mRNA, complete cds.	493	100
578	AAY41716	Homo sapiens	Human PRO860 protein sequence.	5224	100
578	AAB44272	Homo sapiens	Human PRO860 (UNQ421) protein sequence SEQ ID NO:211.	5224	100
578	gi14042832	Homo sapiens	cDNA FLJ14946 fis, clone PLACE2000034, weakly similar to LAR PROTEIN PRECURSOR (EC 3.1.3.48).	3746	93
579	gi7021880	Homo sapiens	cDNA FLJ10054 fis, clone HEMBA1001310.	2306	100
579	gi12653981	Homo sapiens	, TRIAD3 protein, clone MGC:998, mRNA, complete cds.	2306	100
579	gi7109299	Homo sapiens	TRIAD3 mRNA, partial cds.	2013	100
580	gi3288457	Homo sapiens	mRNA for C2 domain containing PI3-kinase.	7615	99 ·
580	gi3059227	Rattus norvegicus	phosphoinositide 3-kinase •	3988	80
580	gi3041786	Mus musculus	Phosphoinositide 3-Kinase-C2gamma	3984	78
581	gi10437125	Homo sapiens	cDNA: FLJ21103 fis, clone CAS04883.	1802	99
581	gi7020867	Homo sapiens	cDNA FLJ20635 fis, clone KAT03466.	786	52
582	gi13937952	Homo sapiens	, Similar to upregulated during skeletal muscle growth 5, clone MGC:14697, mRNA, complete cds.	297	100
582	gi6851054	Rattus norvegicus	DAPIT protein	278	91
582	gi9843791	Mus musculus	stretch regulated skeletal muscle protein	259	84
583	gi7582286	Homo sapiens	BM-007	599	100
583	AAG02907	Homo sapiens	Human secreted protein, SEQ ID NO: 6988.	477	98
583	gi3878572	Caenorhabditis elegans	M01F1.6	161	28
584	gi13477103	Homo sapiens	, clone MGC:1012, mRNA, complete cds.	3001	99
584	gi12052999	Homo sapiens	mRNA; cDNA DKFZp434E1711 (from clone DKFZp434E1711); complete cds.	2619	98
584	gi7020996	Homo sapiens	cDNA FLJ20721 fis, clone HEP15722.	2402	100
585	AAW48892	Homo sapiens	Human guanylate binding protein B (HGBPB).	2645	94
585	gi12803663	Homo sapiens	, guanylate binding protein 1, interferon- inducible, 67kD, clone MGC:3949, mRNA, complete cds.	2000	66
585	gi183002	Homo sapiens	Human guanylate binding protein isoform I (GBP-2) mRNA, complete cds.	2000	66
586	gi7023366	Homo sapiens	cDNA FLJ10983 fis, clone PLACE1001781, weakly similar to PROBABLE PHOSPHOMANNOMUTASE (EC 5.4.2.8).	3218	99
586	gi12052930	Homo sapiens	mRNA; cDNA DKFZp566B1524 (from clone DKFZp566B1524); complete cds.	3216	99
586	gi3395586	Schizosacchar omyces pombe	similarity to phosphomannomutases	1211	43

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
587	gi13537208	Mus musculus	Mel18 and Bmil like ring finger	347	40
587	gi2440074	Homo sapiens	mRNA for RNF3A (DONG1) ring finger protein.	347	37
587	gi13537206	Homo sapiens	hMBLR mRNA, complete cds.	345	40
588	gi14042249	Homo sapiens	cDNA FLJ14610 fis, clone NT2RP1000958, weakly similar to AUTOANTIGEN NGP-1.	2797	99
588	gi14042246	Homo sapiens	cDNA FLJ14608 fis, clone NT2RP1000915, weakly similar to AUTOANTIGEN NGP-1.	2741	99
588	gi6457340	Homo sapiens	E2IG3 (E2IG3) mRNA, complete cds.	2650	100
589	gi7020925 »	Homo sapiens	cDNA FLJ20673 fis, clone KAIA4464.	2232	100
589	gi7682684	Homo sapiens	phosphoprotein associated with GEMs (PAG) mRNA, complete cds.	2222	99
589	gi7707799	Rattus norvegicus	Csk binding protein Cbp	1696	78
590	gi6682873	Homo sapiens	rec mRNA, complete cds.	2002	100
590	gi7230612	Rattus norvegicus	small rec	1916	95
590	gi3881771	Caenorhabditis elegans	contains similarity to Pfam domain: PF01529 (DHHC zinc finger domain), Score=137.4, E-value=8.4e-38, N=1	586	39
591	gi439522	Mus musculus	ribosomal protein S3	678	100
591	gi57728	Rattus rattus	ribosomal protein S3 (AA 1-243)	678	100
591	gi13111933	Homo sapiens	, ribosomal protein S3, clone MGC:3657, mRNA, complete cds.	678	100
592	gi6599070	Homo sapiens	mRNA for LIM domains containing protein 1.	3675	99
592	gi6599307	Mus musculus	LIM domains containing protein 1	2728	76
592	gi13548632	Homo sapiens	partial LIMD1 gene for LIM domains containing 1, exons 1-2, complete sequence.	2690	99
593	gi7020974	Homo sapiens	cDNA FLJ20706 fis, clone KAIA1273.	2824	98
593	gi12082725	Mus musculus	B cell phosphoinositide 3-kinase adaptor	411	29
593	AAG02945	Homo sapiens	Human secreted protein, SEQ ID NO: 7026.	526	100
594	gi11596144	Homo sapiens	STE20-like kinase mRNA, partial cds.	5159	99
594	gi3452473	Rattus norvegicus	serine/threonine protein kinase TAO1	5117	98
594	AAY55937	Homo sapiens	Human SULU3 protein.	4045	100
595	gi695802	Homo sapiens	transcription factor SL1 mRNA, partial cds.	1693	99
595	gi1842206	Mus musculus	TAFI68	1326	76
596	gi7020363	Homo sapiens	cDNA FLJ20335 fis, clone HEP11429.	2940	99
596	AAB65680	Homo sapiens	Novel protein kinase, SEQ ID NO: 208.	2940	99
596	AAB32078	Homo sapiens	Human secreted protein BLAST search protein SEQ ID NO: 136.	826	100
597	gi7020747	Homo sapiens	cDNA FLJ20558 fis, clone KAT11870.	2990	100
597	gi12053175	Homo sapiens	mRNA; cDNA DKFZp434A172 (from clone DKFZp434A172); complete cds.	2990	100
597	gi10439123	Homo sapiens	cDNA: FLJ22650 fis, clone HSI07344.	2166	100
598	gi7023601	Homo sapiens	cDNA FLJ11127 fis, clone PLACE1006225.	1897	100
598	gi12224968	Homo sapiens	mRNA; cDNA DKFZp667E105 (from clone DKFZp667E105).	620	100
598	gi14043433	Homo sapiens	, clone IMAGE:3952677, mRNA, partial cds.	549	41
599	gi6483296	Homo sapiens	CDH9 mRNA for cadherin-9, complete cds.	4132	100

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SEQ ID NO:_	Accession No.	Species	Description	Score	% Identity
599	gi867999	Gallus gallus	chicken cadherin-6B	3044	72
- 599	gi974185	Homo sapiens	mRNA for cadherin-6, complete cds.	3032	72
600	gi5734605	Homo sapiens	mRNA for KARP-1-binding protein 3, complete cds.	750	51
600	gi5734601	Homo sapiens	mRNA for KARP-1-binding protein 1 (KAB1), complete cds.	750	51
600	gi5734603	Homo sapiens	mRNA for KARP-1-binding protein 2 (KAB2), complete cds.	750	51
601	gi10434848	Homo sapiens	cDNA FLJ13028 fis, clone NT2RP3001055, weakly similar to Drosophila melanogaster separation anxiety protein (san) mRNA.	889	100
601	gi10435107	Homo sapiens	cDNA FLJ13194 fis, clone NT2RP3004378, weakly similar to Drosophila melanogaster separation anxiety protein (san) mRNA.	889	100
601	AAB56739	Homo sapiens	Human prostate cancer antigen protein sequence SEQ ID NO:1317.	874	98
602	gi13325182	Homo sapiens	, clone IMAGE:3638994, mRNA, partial cds.	897	100
602	gi12654203	Homo sapiens	, clone IMAGE:3449323, mRNA, partial cds.	560	100
602	gi4514314	Bacillus halodurans	YlqF	260	39
603	gi10954046	Homo sapiens	oxidation protection protein (OXR1) mRNA, complete cds.	1034	97
603	gi13540300	Mus musculus	nucleolar protein C7B	1431	94
603	gi7021988	Homo sapiens	cDNA FLJ10125 fis, clone HEMBA1002954.	1441	99
604	gil150495	Mus musculus	homology to nucleosome assembly proteins; specifically expressed in neurons	211	36
604	gi1161252	Glycine max	nucleosome assembly protein 1	136	40
604	gi5931610	Homo sapiens	mRNA for Nucleosome Assembly Protein 1-like 2, complete cds.	196	37
605	gi7547029	Homo sapiens	GAP-like protein (N61) mRNA, complete cds.	4684	99
605	gi7688683	Homo sapiens	kinesin heavy chain-like protein (KHCHP) mRNA, complete cds.	822	100
605	AAG03378	Homo sapiens	Human secreted protein, SEQ ID NO: 7459.	633	99
606	gi7022593	Homo sapiens	cDNA FLJ10511 fis, clone NT2RP2000656.	1425	100
606	gi12224996	Homo sapiens	mRNA; cDNA DKFZp667G248 (from clone DKFZp667G248).	1031	100
606	gi10436327	Homo sapiens	cDNA FLJ13991 fis, clone Y79AA1002115.	803	100
607	gi8885998	Rattus norvegicus	neuronal C-SRC tyrosine-specific protein kinase	2826	98
607	gi201057	Mus musculus	tyrosine-specific protein kinase	2822	98
607	gi338460	Homo sapiens	Human c-src-1 proto-oncogene, exon 12.	2815	98
608	gi7243633	Homo sapiens	RB-associated KRAB repressor (RBAK) mRNA, complete cds.	3993	100
608	gi7243635	Mus musculus	RB-associated KRAB repressor	3025	78
608	gi10434235	Homo sapiens	cDNA FLJ12629 fis, clone NT2RM4001828, moderately similar to ZINC FINGER PROTEIN 84.	1881	73
609	gi7008402	Homo sapiens	kappa B-ras I mRNA, complete cds.	982	100
609	gi14042659	Homo sapiens	cDNA FLJ14843 fis, clone	978	99

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
			PLACE1000040, weakly similar to TRANSFORMING PROTEIN P21/K- RAS 2B.		
609	gi7239257	Mus musculus	kappaB-Ras l	952	94
610	gi13625164	Homo sapiens	ankyrin mRNA, complete cds.	426	100
610	gi12698638	Homo sapiens	ankyrin-repeat family A protein 2 (ANKRA2) mRNA, complete cds.	426	100
610	gi10434525	Homo sapiens	cDNA FLJ12814 fis, clone NT2RP2002520, weakly similar to Homo sapiens transcription factor RFX-B (RFXB) mRNA.	426	100
611	gi7959841	Homo sapiens	PRO1853	510	100
611	AAG01282	Homo sapiens	Human secreted protein, SEQ ID NO: 5363.	301	100
612	gi5757703	Mus musculus	syntrophin-associated serine-threonine protein kinase	7464	92
612	gi13537204	Homo sapiens	mRNA for MAST205, complete cds.	4616	68
612	gi406058	Mus musculus	protein kinase	4569	65
613	gi7020724	Homo sapiens	cDNA FLJ20545 fis, clone KAT11476.	1780	100
613	AAB63186	Homo sapiens	Human secreted protein sequence encoded by gene 3 SEQ ID NO:112.	1693	100
613	gi7243701	Drosophila melanogaster	WDS	1574	91
614	gi13383476	Homo sapiens	NUB1 (NUB1) mRNA, complete cds.	3109	100
614	gi5360093	Homo sapiens	NY-REN-18 antigen mRNA, complete cds.	2958	95
614	gi863014	Mus musculus	BS4 peptide	2671 .	84
615	AAB87345	Homo sapiens	Human gene 4 encoded secreted protein HDPFY41, SEQ ID NO:86.	4534	100
615	gi4886489	Homo sapiens	mRNA; cDNA DKFZp564L2123 (from clone DKFZp564L2123); partial cds.	2892	99
615	gi12711793	Homo sapiens	estrogen regulated LIV-1 protein (LIV-1) mRNA, complete cds.	1171	39
616	gi7638247	Homo sapiens	mesenchymal stem cell protein DSCD75 mRNA, complete cds.	1063	100
616	gi12654929	Homo sapiens	, mesenchymal stem cell protein DSCD75, clone MGC:5515, mRNA, complete cds.	1063	100
616	AAB03956	Homo sapiens	Human mesenchymal stem cell polypeptide.	1063	100
617	gi7582304	Homo sapiens	BM-016	584	100
617	AAW78199	Homo sapiens	Human secreted protein encoded by gene 74 clone HGBAC11.	562	98
617	AAW85610	Homo sapiens	Secreted protein clone eh80_1.	562	98
618	gi13603398	Homo sapiens	mRNA for SEZ6L, complete cds.	4199	98
618	gi13185723	Homo sapiens	n 1755 can be A, G, C, or T	2164	49
618	AAB70537	Homo sapiens	Human PRO7 protein sequence SEQ ID NO:14.	2164	49
619	gi3880445	Caenorhabditis elegans	contains similarity to Pfam domain: PF02214 (K+ channel tetramerisation domain), Score=79.5, E-value=2.3e-20, N=1	327	40
619	AAY34129	Homo sapiens.	Human potassium channel K+Hnov28.	195	40
619	AAZ11907_ aa1	Homo sapiens	Human potassium channel K+Hnov28 cDNA (5' splice variant 1).	195	40
620	gi10437116	Homo sapiens	cDNA: FLJ21097 fis, clone CAS03931.	1146	100
620	gi14250732	Homo sapiens	, chromosome 11 open reading frame 14, clone MGC:12847, mRNA, complete cds.	1146	100
620	gi13276621	Homo sapiens	mRNA; cDNA DKFZp761G1913 (from	378	43

SEQ ID	Accession	Species	Description	Score	%
NO:	No.	SP10.03			Identity
			clone DKFZp761G1913).		
621	gi10437078	Homo sapiens	cDNA: FLJ21069 fis, clone CAS01594.	955	58
621	gi5911935	Homo sapiens	mRNA; cDNA DKFZp586N1922 (from clone DKFZp586N1922); partial cds.	867	100
621	AAB27870	Homo sapiens	Protein fragment encoded by gene 27.	657 ·	100
622	gi13097159	Homo sapiens	, tumor protein, translationally-controlled 1, clone MGC:5308, mRNA, complete cds.	898	100
622	gi14043771	Homo sapiens	, clone MGC:14243, mRNA, complete cds.	898	100
622	gi7573519	Homo sapiens	TPT1 gene for translationally controlled tumor protein (TCTP), exons 1-6.	898	100
623	gi7020339 -	Homo sapiens	cDNA FLJ20320 fis, clone HEP08923.	1135	100
623	AAB18972	Homo sapiens	Amino acid sequence of a human transmembrane protein.	1135	100
623	gi1314162	Schizosacchar omyces pombe	seven transmembrane protein	217	29
624	gi6467990	Mus musculus	PDZ domain actin binding protein Shroom	4816	66
624	gi6467992	Mus musculus	actin binding protein ShroomS	4816	66
624	gi13938323	Homo sapiens	, Similar to shroom, clone IMAGE:3349317, mRNA, partial cds.	4006	99
625	gi12804029	Homo sapiens	, clone IMAGE:3940519, mRNA, partial cds.	1551	100
625	AAY21850	Homo sapiens	Human signal peptide-contianing protein (SIGP) (clone ID 1880830).	1109	100
625	gi8655657	Homo sapiens	mRNA; cDNA DKFZp762O076 (from clone DKFZp762O076).	593	57
626	gi7328140	Homo sapiens	mRNA; cDNA DKFZp762D096 (from clone DKFZp762D096); partial cds.	601	100
626	gi13436341	Homo sapiens	, Similar to RIKEN cDNA 1600014C10 gene, clone MGC:10922, mRNA, complete cds.	384	100
627	gi1293559	Mus musculus	astrotactin	4312	95
627	gi6502571	Mus musculus	astrotactin2	2580	51
627	gi6502573	Homo sapiens	astrotactin2 (ASTN2) mRNA, complete cds.	2569	51
628	AAY73387	Homo sapiens	HTRM clone 3340290 protein sequence.	1439	95
628	AAY48312	Homo sapiens	Human prostate cancer-associated protein 9.	1073	84
628	gi12654077	Homo sapiens	, clone IMAGE:3458173, mRNA, partial cds.	1045	86
629	gi11095188	Homo sapiens	dipeptidyl peptidase 8 (DPP8) mRNA, complete cds.	3521	99
629	gi14042790	Homo sapiens	cDNA FLJ14920 fis, clone PLACE1007416, weakly similar to DIPEPTIDYL PEPTIDASE IV (EC 3.4.14.5).	2457	99
629	gi7020273	Homo sapiens	cDNA FLJ20283 fis, clone HEP04088.	2483	100
630	gi11095188	Homo sapiens	dipeptidyl peptidase 8 (DPP8) mRNA, complete cds.	2560	99
630	gi14042790	Homo sapiens	cDNA FLJ14920 fis, clone PLACE1007416, weakly similar to DIPEPTIDYL PEPTIDASE IV (EC 3.4.14.5).	2457	99
630	gi11095192	Homo sapiens	dipeptidyl peptidase 8 (DPP8) mRNA, partial cds, alternatively spliced.	2482	100
631	gi7020611	Homo sapiens	cDNA FLJ20481 fis, clone KAT07534.	2211	99
631	AAY57908	Homo sapiens	Human transmembrane protein HTMPN-	975	44

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
NO:	No.		32.		lucillity
631	AAB54284	Homo sapiens	Human pancreatic cancer antigen protein sequence SEQ ID NO:736.	516	40
632	gi35700	Homo sapiens	Human mRNA for phosphoriobosyl pyrophosphate synthetase subunit II (EC 2.7.6.1).	1596	99
632	gi206434	Rattus norvegicus	phosphoribosyl pyrophosphate synthetase II	1585	98
632	gi56979	Rattus norvegicus	ribose-phosphate pyrophosphokinase subunit II (AA 1-318)	1585	98
633	gi11181620	Homo sapiens	Rag D mRNA, complete cds.	1276	100
633	gi6808148	Homo sapiens	mRNA; cDNA DKFZp761H171 (from clone DKFZp761H171); partial cds.	1276	100
633	AAB56443	Homo sapiens	Human prostate cancer antigen protein sequence SEQ ID NO:1021.	1276	100
634	gi6807893	Homo sapiens	mRNA; cDNA DKFZp434H2226 (from clone DKFZp434H2226); partial cds.	1079	100
635	gi10435042	Homo sapiens	cDNA FLJ13152 fis, clone NT2RP3003385, highly similar to Mus musculus SKD3 mRNA.	3495	100
635	gi4958935	Rattus norvegicus	suppressor of potassium transport defect 3	3085	88
635	gi563129	Mus musculus	SKD3	3066	88
636	AAB20322	Homo sapiens	Human protein phosphatase and kinase protein-1.	1770	100
636	gi1903458	Dictyostelium discoideum	myosin heavy chain kinase B	236	32
636	gi2104701	Mus musculus	elongation factor-2 kinase	199	29
637	gi7670003	Homo sapiens	mRNA; cDNA DKFZp434P0531 (from clone DKFZp434P0531).	1850	100
637	gi7417474	Homo sapiens	chromosome 14 clone RP11-493G17 and CTD-2516D11 map 14q24.3, complete sequence.	1251 .	49
637	gi7018538	Homo sapiens	mRNA; cDNA DKFZp434P0111 (from clone DKFZp434P0111); partial cds.	330	43
638	gi7022367	Homo sapiens	cDNA FLJ10375 fis, clone NT2RM2001950.	3056	100
638	AAY53026	Homo sapiens	Human secreted protein clone cn922_5 protein sequence SEQ ID NO:58.	1752	95
638	gi4336692	Drosophila melanogaster	Abnormal X segregation	816	37
639	gi7020972	Homo sapiens	cDNA FLJ20705 fis, clone KAIA1571.	3641	99
639	gi12007334	Homo sapiens	IRS-1 PH domain binding protein PHIP mRNA, complete cds.	3632	99
639	gi14286226	Homo sapiens	, pleckstrin homology domain interacting protein, clone MGC:15187, mRNA, complete cds.	3632	99
640	gi7689025	Homo sapiens	uncharacterized hypothalamus protein HT013 mRNA, complete cds.	978	96
641	gi9937505	Homo sapiens	PLIC-2 mRNA, complete cds.	3167	100
641	gi6563288	Homo sapiens	ubiquitin-like product Chap1/Dsk2 mRNA, complete cds.	3162	99
641	AAB47122	Homo sapiens	Human Chap1.	3162	99
642	AAY53001	Homo sapiens	Human secreted protein clone dn834_1 protein sequence SEQ ID NO:8.	811	100
642	AAG01114	Homo sapiens	Human secreted protein, SEQ ID NO: 5195.	641	99
642	gi12652989	Homo sapiens	, clone MGC:2495, mRNA, complete cds.	489	57
643	gi7021064	Homo sapiens	cDNA FLJ20761 fis, clone HEP00317.	2240	100

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
643	gi10438264	Homo sapiens	cDNA: FLJ22019 fis, clone HEP07982.	2187	98
643	gi577428	Rattus	Ca2+-dependent activator protein;	1787	77
		norvegicus	calcium-dependent actin-binding protein		
644	gi7023651	Homo sapiens	cDNA FLJ11159 fis, clone PLACE1006966.	2865	99
644	gi7023118	Homo sapiens	cDNA FLJ10835 fis, clone NT2RP4001210.	1253	100
644	gi600058	Saccharomyce s cerevisiae	N1342	710	39
645	gi7020012	Homo sapiens	cDNA FLJ20121 fis, clone COL05942.	1334	99
646	gi14336697	Homo sapiens	16p13.3 sequence section 2 of 8.	609	100
646	gi13436122	Homo sapiens	, non-metastatic cells 4, protein expressed in, clone MGC:11088, mRNA, complete cds.	609	100
646	gi1945762	Homo sapiens	H.sapiens mRNA for nucleoside- diphosphate kinase.	609	100
647	AAB24225	Homo sapiens	Human vesicle associated protein 4 SEQ ID NO:4.	2946	99
647	gi10439139	Homo sapiens	cDNA: FLJ22662 fis, clone HSI08080.	2703	99
647	AAB58427	Homo sapiens	Lung cancer associated polypeptide sequence SEQ ID 765.	1711	99
648	gi7020604	Homo sapiens	cDNA FLJ20477 fis, clone KAT07271.	2639	99
648	gi6672090	Drosophila melanogaster	Vegetable	578	32
649	gi12802986	Homo sapiens	, ring finger protein 24, clone MGC:1815, mRNA, complete cds.	811	100
649	gi5420200	Homo sapiens	Novel human mRNA from chromosome 20, similar to SW:GOLI_DROME Q06003 GOLIATH PROTEIN.	811	100
649	gi5102892	Homo sapiens	mRNA full length insert cDNA clone EUROIMAGE 566628.	566	100
650	gi6841346	Homo sapiens	HSPC054	497	98
651	gi7209305	Homo sapiens	mRNA for FLJ00002 protein, partial cds.	7637	100
651	gi6599226	Homo sapiens	mRNA; cDNA DKFZp434L0827 (from clone DKFZp434L0827); partial cds.	3519	100
651	gi10440406	Homo sapiens	mRNA for FLJ00036 protein, partial cds.	3457	99
652	gi7018505	Homo sapiens	mRNA; cDNA DKFZp434E2220 (from clone DKFZp434E2220).	2470	100
652	gi14042579	Homo sapiens	cDNA FLJ14796 fis, clone NT2RP4001235.	2466	99
652	gi7018507	Homo sapiens	mRNA; cDNA DKFZp434O0420 (from clone DKFZp434O0420).	2466	99
653	gi552196	Plasmodium lophurae	histidine-rich protein	192	40
653	gi160362	Plasmodium falciparum	knob protein	178	42
653	gi3845095	Plasmodium falciparum	knob-associated His-rich protein	172	40
654	AAY70539	Homo sapiens	Human Factor 8 Homologue.	1353	83
654	gi14043498	Homo sapiens	, Similar to neuropilin 1, clone MGC:12920, mRNA, complete cds.	189	34
654	gi7271465	Homo sapiens	soluble neuropilin-1 mRNA, complete cds.	189	34
655	gi7019959	Homo sapiens	cDNA FLJ20087 fis, clone COL03793.	3964	100
655	gi13569705	Homo sapiens	channel kinase 2 (CHAK2) mRNA, complete cds.	3942	99
655	AAY95433	Homo sapiens	Human calcium channel SOC-2/CRAC-1 C-terminal polypeptide.	1172	71
656	gi6094668	Homo sapiens	BAC clone RP11-343N14 from 2,	208	100

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
			complete sequence.		
656	gi10435833	Homo sapiens	cDNA FLJ13729 fis, clone PLACE3000121, weakly similar to VESICULAR TRAFFIC CONTROL PROTEIN SEC15.	208	100
656	gi2827162	Rattus norvegicus	rsec15	160	73
657	gi10434153	Homo sapiens	cDNA FLJ12580 fis, clone NT2RM4001116, weakly similar to HYPOTHETICAL 216.3 KD PROTEIN R06F6.8 IN CHROMOSOME II.	1806	99
657	gi12053255	Homo sapiens	mRNA; cDNA DKFZp434D105 (from clone DKFZp434D105); complete cds.	1806	99
657	gi5901808	Drosophila melanogaster	BcDNA.GH03694	619	56
658	gil1181618	Homo sapiens	Rag C mRNA, complete cds.	2072	100
658	gi12007486	Homo sapiens	GTPase-interacting protein 2 mRNA, complete cds.	2069	99
658	gi13529335	Mus musculus	Similar to Rag C protein	2039	98
659	gi13537208	Mus musculus	Mel18 and Bmil like ring finger	347	40
659	gi2440074	Homo sapiens	mRNA for RNF3A (DONG1) ring finger protein.	347	37
659	gi13537206	Homo sapiens	hMBLR mRNA, complete cds.	345	40
660	gi7023690	Homo sapiens	cDNA FLJ11184 fis, clone PLACE1007507.	1043	99
661	gi7020878	Homo sapiens	cDNA FLJ20641 fis, clone KAT02782.	2552	99
661	gi11992034	Rattus norvegicus	antisense RNA overlapping MCH protein	1609	65
662	AAB56646	Homo sapiens	Human prostate cancer antigen protein sequence SEQ ID NO:1224.	915	98
662	gi12053357	Homo sapiens	mRNA; cDNA DKFZp586G2122 (from clone DKFZp586G2122); complete cds.	900	100
662	AAB36598	Homo sapiens	Human FLEXHT-20 protein sequence SEQ ID NO:20.	791	59
663	AAW93947	Homo sapiens	Human regulatory molecule HRM-3 protein.	1732	100
663	gi3288459	Homo sapiens	mRNA for transcription elongation factor TFIIS.h.	1673	100
663	gi3288547	Mus musculus	transcription elongation factor TFIIS.h	1543	90
664	gi14042893	Homo sapiens	cDNA FLJ14984 fis, clone Y79AA1000349, highly similar to M.musculus Spnr mRNA for RNA binding protein.	3478	100
664	gi13377630	Homo sapiens	spermatid perinuclear RNA-binding protein mRNA, complete cds.	3459	99
664	gi12053237	Homo sapiens	mRNA; cDNA DKFZp434N214 (from clone DKFZp434N214); complete cds.	3406	100
665	gi10436573	Homo sapiens	cDNA FLJ14183 fis, clone NT2RP2004920, weakly similar to TRANSCRIPTIONAL REGULATOR ATRX.	4423	99
665	gi10434345	Homo sapiens	cDNA FLJ12693 fis, clone NT2RP1000324.	4369	99
665	AAB27235	Homo sapiens	Human EXMAD-13 SEQ ID NO: 13.	3346	100
666	gi9858154	Homo sapiens	tubby super-family protein (TUSP) mRNA, complete cds, alternatively spliced.	3598	100
666	gi9502082	Homo sapiens	tubby super-family protein (TUSP) mRNA, complete cds.	3556	100

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
666	gi9502080	Mus musculus	tubby super-family protein	3505	98
667	gi7106796	Homo sapiens	HSPC203	554	100
667	gi9963859	Homo sapiens	PTD019 mRNA, complete cds.	554	100
667	AAY35987	Homo sapiens	Extended human secreted protein sequence, SEQ ID NO. 236.	554	100
668	gi6996442	Homo sapiens	CTL1 gene.	3398	99
668	gi6996589	Rathis norvegicus	CTL1 protein	3291	96
668	gi6996587	Torpedo marmorata	CTL1 protein	2454	71
669	gi6808165	Homo sapiens	mRNA; cDNA DKFZp761A052 (from clone DKFZp761A052).	2265	100
669	gi10439058	Homo sapiens	cDNA: FLJ22607 fis, clone HSI04846.	1992	100
669	gi7673616	Mus musculus	DXImx46e protein	1958	98
670	gi6808252	Homo sapiens	mRNA; cDNA DKFZp434D1319 (from clone DKFZp434D1319); partial cds.	2336	100
670	gi170035	Glycine max	N-75	221	27
670	gi18576	Glycine max	pre-pro polypeptide (AA -25 to 284)	219	27
671	AAW93947	Homo sapiens	Human regulatory molecule HRM-3 protein.	1116	99
671	gi3288459	Homo sapiens	mRNA for transcription elongation factor TFUS.h.	1057	99
671	gi3288547	Mus musculus	transcription elongation factor TFIIS.h	950	86
672	gi10434615	Homo sapiens	cDNA FLJ12875 fis, clone NT2RP2003777.	1818	99
672	gi8778741	Arabidopsis thaliana	T30E16.12	254	27
672	gi6520214	Arabidopsis thaliana	ZCF61	228	29
673	AAB88424	Homo sapiens	Human membrane or secretory protein clone PSEC0197.	3032	99
673	gi9294464	Arabidopsis thaliana	long-chain-fatty-acid-CoA ligase-like protein	581	37
673	gi699196	Mycobacteriu m leprae	4-coumarate-coA ligase	326	45
674	gi7022969	Homo sapiens	cDNA FLJ10747 fis, clone NT2RP3001799.	3378	99
674	AAY86211	Homo sapiens	Nuclear transport protein clone hfb066 protein sequence.	1432	87
674	gi10439560	Homo sapiens	cDNA: FLJ23007 fis, clone LNG00451.	703	100
675	gi7021968	Homo sapiens	cDNA FLJ10111 fis, clone HEMBA1002696.	2753	99
675	gi14017768	Mus musculus	FLJ10111	2214	92
675	gi10440211	Homo sapiens	cDNA: FLJ23501 fis, clone LNG02837.	2160	90
676	gi7021968	Homo sapiens	cDNA FLJ10111 fis, clone HEMBA1002696.	2728	98
676	gi14017768	Mus musculus	FLJ10111	2200	90
676	gi10440211	Homo sapiens	cDNA: FLJ23501 fis, clone LNG02837.	2237	92
677	gi7019869	Homo sapiens	cDNA FLJ20036 fis, clone COL00219.	2834	100
677	gi12723779	Lactococcus lactis subsp. lactis	UNKNOWN PROTEIN	306	35
677	gi8885520	Streptococcus gordonii	streptococcal hemagglutinin	297	29
678	gi10437508	Homo sapiens	cDNA: FLJ21415 fis, clone COL04030.	1129	100
679	gi3135314	Homo sapiens	chromosome 7q22 sequence, complete sequence.	1226	100
679	gi6752287	Homo sapiens	Novel human gene mapping to chomosome X.	390	43

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
679	AAB28327	Homo sapiens	Human secreted protein BLAST search protein SEQ ID NO: 111.	265	100
680	gi3135314	Homo sapiens	chromosome 7q22 sequence, complete sequence.	1199	95
680	gi6752287	Homo sapiens	Novel human gene mapping to chomosome X.	363	41
680	AAB28327	Homo sapiens	Human secreted protein BLAST search protein SEQ ID NO: 111.	265	100
681	gi10439089	Homo sapiens	cDNA: FLJ22626 fis, clone HSI06109.	2120	99
681	gi11044557	Homo sapiens	Human DNA sequence from clone RP11- 42415 on chromosome 6 Contains a novel gene, STSs, GSSs and a CpG island, complete sequence.	. 1185	64
681	gi12654241	Homo sapiens	, Similar to splicing factor, arginine/serine-rich 4 (SRp75), clone MGC:5283, mRNA, complete cds.	949	98
682	gi14042277	Homo sapiens	cDNA FLJ14626 fis, clone NT2RP2000288.	3029	99
682	gi7022410	Homo sapiens	cDNA FLJ10402 fis, clone NT2RM4000457.	2279	100
682	gi6841196	Homo sapiens	HSPC273	1086	100
683	gi2815604	Homo sapiens	Opa-interacting protein OIP2 mRNA, partial cds.	1364	100
683	AAB63276	Homo sapiens	Human breast cancer associated antigen protein sequence SEQ ID NO:638.	839	96
683	AAB63406	Homo sapiens	Human breast cancer associated antigen protein sequence SEQ ID NO:768.	839	96
684	AAB07228	Homo sapiens	Human prostate cancer predisposing protein HPC2.	4325	100
684	AAY99850	Homo sapiens	Human sulphatase G.	4315	99
684	gi10946497	Pan troglodytes	ELAC2	4283	98
685	gi7688979	Homo sapiens	uncharacterized bone marrow protein BM042	895	100
685 .	AAB36580	Homo sapiens	Human FLEXHT-2 protein sequence SEQ ID NO:2.	895	100
685	AAB34771	Homo sapiens	Human secreted protein fragment encoded by DNA clone vq23 1.	888	99
686	gi10438990	Homo sapiens	cDNA: FLJ22559 fis, clone HSI01591.	1897	100
686	gi8954034	Arabidopsis thaliana	F10K1.17	162	31
687	gi7020674	Homo sapiens	cDNA FLJ20515 fis, clone KAT09889.	2027	100
687	AAB20331	Homo sapiens	Human protein phosphatase and kinase protein-10.	1472	92
687	AAB73226	Homo sapiens	Human phosphatase NP 060746 h.	576 1019	100
688	gi6688145	Homo sapiens	mRNA for NICE-3 protein, clone 1023j12.		
688	gi4689120	Homo sapiens	HSPC012	717	93
688	gi12655055	Homo sapiens	, DKFZP586G1722 protein, clone MGC:1147, mRNA, complete cds.	717	93
689	gi7023701	Homo sapiens	cDNA FLJ11190 fis, clone PLACE1007583.	1317	100
690	gi6469703	Mycobacteriu m tuberculosis	DipZ	203	31
691	gi13676779	Mus musculus		1939	93
691	gi13752369	Gallus gallus	ring finger protein	1888	91
691	gi13752371	Xenopus laevis	ring finger-H2 protein	1537	76
692	gi458255	Homo sapiens	Human X-linked PEST-containing	2849	99

SEQ ID	Accession	Species	Description	Score	%
<u>NO:</u>	No.				Identity
	-:460047	Homo sapiens	transporter (XPCT) gene, exon 6. Human X-linked PEST-containing	2766	99
692	gi458247	Homo sapiens	transporter (XPCT) mRNA, partial cds.	2700	33
692	gi2944356	Mus musculus	X-linked PEST-containing transporter	2249	88
693	gi14042736	Homo sapiens	cDNA FLJ14888 fis, clone	2034	99
			PLACE1003762.		
693	gi6841178	Homo sapiens	HSPC264	2019	99
694	gi7023413	Homo sapiens	cDNA FLJ11012 fis, clone	2377	99
			PLACE1003190, weakly similar to SOFI PROTEIN.		
694	gi14042745	Homo sapiens	cDNA FLJ14893 fis, clone	2377	99
		٠	PLACE1004302, weakly similar to SOF1 PROTEIN.		
694	gi5912184	Homo sapiens	mRNA; cDNA DKFZp564O0463 (from clone DKFZp564O0463); partial cds.	1159	99
695	gi7022931	Homo sapiens	cDNA FLJ10724 fis, clone NT2RP3001176.	2683	99
695	gi14198202	Mus musculus	Similar to melanoma antigen recognized by T cells 2	2126	82
695	gi4826524	Homo sapiens	Novel human gene mapping to chomosome 1.	982	92
696	gi7022990	Homo sapiens	cDNA FLJ10761 fis, clone	2119	99
0,00	8		NT2RP3004669, weakly similar to		
	ì		ETHANOLAMINE KINASE (EC		
	-:0000050	Illama amiana	2.7.1.82). ethanolamine kinase (EKII) mRNA,	930	56
696	gi9998952	Homo sapiens	complete cds.	930	. 36
696	gi532128	Drosophila melanogaster	ethanolamine kinase	525	45
697	gi186774	Homo sapiens	Human Kruppel related zinc finger protein (HTF10) mRNA, complete cds.	986	38
697	gi5441615	Canis familiaris	zinc finger protein	988	37
697	gi38032	Homo sapiens	Human ZNF43 mRNA.	947	36
698	gi13537202	Homo sapiens	PC-LKC mRNA for protocadherin LKC, complete cds.	2877	100
698 .	gi7020017	Homo sapiens	cDNA FLJ20124 fis, clone COL06056.	2862	99
698	AAY01410	Homo sapiens	Secreted protein encoded by gene 28 clone HE9ND43.	963	100
699	gi7688977	Homo sapiens	uncharacterized bone marrow protein BM041	888	100
699	AAY86515	Homo sapiens	Human gene 71-encoded protein fragment, SEQ ID NO:430.	888	100
699	gi7018421	Homo sapiens	mRNA; cDNA DKFZp564J157 (from clone DKFZp564J157).	880	99
700	gi7209307	Homo sapiens	mRNA for FLJ00003 protein, partial cds.	1102	100
700	gi14276857	Homo sapiens	PC2-glutamine-rich-associated protein (PCOAP) mRNA, complete cds.	429	93
700	gi14043091	Homo sapiens	, clone IMAGE:3350171, mRNA, partial cds.	429	93
701	gi7020678	Homo sapiens	cDNA FLJ20517 fis, clone KAT10235.	2821	99
701	gi10177966	Arabidopsis thaliana	uridine kinase-like protein	1068	44
701	gi496728	Saccharomyce s cerevisiae	uridine kinase	775	37
702	gi7022789	Homo sapiens	cDNA FLJ10634 fis, clone	1512	100
			NT2RP2005654, weakly similar to		
702	A A D 67446	II	CYSTEINE STRING PROTEIN.	1512	100
702	AAB67446	Homo sapiens	Amino acid sequence of a human	1512	100

SEQ ID	Accession	Species	Description	Score	%
NO:	No.				Identity
702	A A CO1052	**	chaperone polypeptide.	<u> </u>	ļ
	AAG01952	Homo sapiens	Human secreted protein, SEQ ID NO: 6033.	422	98
703	gi7021321	Homo sapiens	Gemin4 mRNA, complete cds.	5481	99
703	gi10945430	Homo sapiens	chromosome 17 clone PAC P579 HC90, HC71AC, HC6 and HC56 genes, complete sequence.	5452	100
703	gi7018412	Homo sapiens	mRNA; cDNA DKFZp434D174 (from clone DKFZp434D174).	4359	99
704	gi9964287	Homo sapiens	hypertension-related calcium-regulated gene mRNA, complete cds.	1129	100
704	gi10434820	Homo sapiens	cDNA FLJ13008 fis, clone NT2RP3000456.	1129	100
704	gi12803673	Homo sapiens	, HT002 protein; hypertension-related calcium-regulated gene, clone MGC:3418, mRNA, complete cds.	1129	100
705	gi10435947	Homo sapiens	cDNA FLJ13814 fis, clone THYRO1000368.	3588	99
705	gi3878402	Caenorhabditis elegans	similar to C2 domain	300	25
705	gi3002479	Leishmania major	L3162.1	198	25
706	gi11907998	Homo sapiens	BCL-6 corepressor (BCOR) mRNA, complete cds; alternatively spliced.	2449	100
706	gi7020277	Homo sapiens	cDNA FLJ20285 fis, clone HEP04260.	1131	99
706	gi10432606	Homo sapiens	cDNA FLJ11362 fis, clone HEMBA1000244.	458	50
707	gi7768662	Homo sapiens	C4ST mRNA for chondroitin 4- sulfotranseferase, complete cds.	1870	100
707	gi8925966	Homo sapiens	chondroitin 4-O-sulfotransferase 1 mRNA, complete cds.	1870	100
707	gi7572958	Homo sapiens	mRNA for chondroitin-4-sulfotransferase (C4ST gene).	1865	99
708	gi2731561	Homo sapiens	ATP receptor subunit (P2X5) mRNA, complete cds.	2167	96
708	gi1552522	Homo sapiens	Human ionotropic ATP receptor P2X5a mRNA, complete cds.	2131	96
708	gi3387944	Homo sapiens	clone 24793 ionotropic ATP receptor P2X5b mRNA, complete cds.	1608	99
709	gi7021105	Homo sapiens	cDNA FLJ20793 fis, clone COL00343.	1587	100
709	gi7206854	Caenorhabditis elegans	contains similarity to Pfam family PF00085 (Thioredoxins), Score 113, E=9.6e-33, N=1	435	29
709	gi13775331	Caenorhabditis elegans	contains similarity to Pfam family PF00085 (Thioredoxin), score=320.7, E=1.8e-95, N=3	297	28
710	AAY04315	Homo sapiens	Human secreted protein encoded by gene 23.	385	100
710	AAB12155	Homo sapiens	Hydrophobic domain protein isolated from HT-1080 cells.	385	100
711	gi13624098	Homo sapiens	cervical cancer 1 protooncogene protein p40 mRNA, complete cds.	520	100
711	gi12653253	Homo sapiens	, DKFZP586A011 protein, clone MGC:8483, mRNA, complete cds.	520	100
711	gi4886473	Homo sapiens	mRNA; cDNA DKFZp586A011 (from clone DKFZp586A011); partial cds.	520	100
712	gi927415	Homo sapiens	H.sapiens mRNA for carnitine acetyltransferase.	3209	98
712	gi13879380	Mus musculus	Similar to carnitine acetyltransferase	3010	90

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
712	gi758632	Mus musculus	carnitine acetyltransferase	2967	89
713	gi9437507	Homo sapiens	TERA	1198	100
713	gi10439906	Homo sapiens	cDNA: FLJ23279 fis, clone HEP06870.	1198	100
713	gi12652565	Homo sapiens	, TERA protein, clone MGC:1093, mRNA, complete cds.	1198	100
714	gi7023336	Homo sapiens	cDNA FLJ10964 fis, clone PLACE1000748.	1196	100
714	gi14198104	Homo sapiens	, clone MGC:16981, mRNA, complete cds.	1196	100
714	gi7023823	Homo sapiens	cDNA FLJ11269 fis, clone PLACE1009190.	661	100
715	AAB67579	Homo sapiens	Amino acid sequence of a human hydrolytic enzyme HYENZ11.	2740	100
715	gi7020019	Homo sapiens	cDNA FLJ20125 fis, clone COL06152.	1973	99
715	gi13527857	Drosophila melanogaster	pol polyprotein	298	26
716	gi2218077	Homo sapiens	gravin mRNA, complete cds.	8920	99
716	AAW53863	Homo sapiens	Human gravin polypeptide.	8868	99
716	AAB15380	Homo sapiens	Human gravin protein sequence.	8868	99
717	gi7021891	Homo sapiens	cDNA FLJ10060 fis, clone HEMBA1001407.	2306	99
717	gi10433215	Homo sapiens	cDNA FLJ11856 fis, clone HEMBA1006789.	1959	86
717	gi14042890	Homo sapiens	cDNA FLJ14982 fis, clone Y79AA1000258.	1959	86
718	gi6224691	Homo sapiens	Na+/sulfate cotransporter SUT-1 (SUT-1) mRNA, complete cds.	3271	100
718	AAB36158	Homo sapiens	Novel human transporter protein SEQ ID NO: 2.	3268	99
718	AAB23625	Homo sapiens	Human secreted protein SEQ ID NO: 50.	3268	99
719	gi7020123	Homo sapiens	cDNA FLJ20189 fis, clone COLF0657.	1264	99
719	gi14328904	Homo sapiens	fetal globin-inducing factor (FGIF) mRNA, complete cds.	1262	99
719	AAB71861	Homo sapiens	Human FGIF.	1262	99
720	gi6690250	Homo sapiens	clone HQ0659 PRO0659 mRNA, complete cds.	926 .	100
720	gi12654109	Homo sapiens	, PRO0659 protein, clone MGC:4888, mRNA, complete cds.	926	100
721	gi608025	Homo sapiens	Human ankyrin G (ANK-3) mRNA, complete cds.	580	32
721	gi3885972	Rattus norvegicus	270 kDa ankyrin G isoform	575	32
721	gi178646	Homo sapiens	Human erythroid ankyrin mRNA, complete cds.	609	35
722	gi7020915	Homo sapiens	cDNA FLJ20666 fis, clone KA1A608.	1229	96
722	gi3169096	Schizosacchar omyces pombe	possible pre-mRNA processing by similarity to yeast prp39	420	37
722	gi1458279	Caenorhabditis elegans	contains similarity to TPR domains	252	29
723	gi7020729	Homo sapiens	cDNA FLJ20548 fis, clone KAT11542.	2200	100
723	gi10434720	Homo sapiens	cDNA FLJ12942 fis, clone NT2RP2005139, weakly similar to 2-5A- DEPENDENT RIBONUCLEASE (EC 3.1.26).	2200	100
723	gi11967781	Homo sapiens	ANKRD2 gene for skeletal muscle ankyrin repeat, exons 1-9.	174	30
724	gi10433458	Homo sapiens	cDNA FLJ12068 fis, clone HEMBB1002329.	2903	99
724	gi10434339	Homo sapiens	cDNA FLJ12690 fis, clone	2898	99

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
			NT2RM4002567.		
724	gi10436665	Homo sapiens	cDNA FLJ14252 fis, clone OVARC1001341.	2167	99
725	gi10434638	Homo sapiens	cDNA FLJ12889 fis, clone NT2RP2004098, weakly similar to ADENYLATE CYCLASE (EC 4.6.1.1).	3026	100
725	gi14250313	Homo sapiens	, clone MGC:16864, mRNA, complete cds.	3026	100
725	gi7020356	Homo sapiens	cDNA FLJ20331 fis, clone HEP10410.	1914	99
726	AAY13947	Homo sapiens	Human transmembrane protein, HP10495.	655	100
726	AAY07878	Homo sapiens	Human secreted protein fragment encoded from gene 27.	655	100
726	gi6841296	Homo sapiens	HSPC323	449	85
727	gi7159733	Homo sapiens	mRNA for ETAA16 protein.	4318	100
727	AAB10622	Homo sapiens	Human Ewing tumor associated antigen protein.	4318	100
728	gi7020138	Homo sapiens	cDNA FLJ20199 fis, clone COLF1162.	2123	99
728	AAY91948	Homo sapiens	Human cytoskeleton associated protein 3 (CYSKP-3).	1650	99
728	gi7020210	Homo sapiens	cDNA FLJ20246 fis, clone COLF6458.	1474	99
729	gi13182775	Homo sapiens	CDA11 mRNA, complete cds.	1495	99
729	gi13937914	Homo sapiens	, clone MGC:12519, mRNA, complete cds.	973	97
729	gi2257524	Schizosacchar omyces pombe	HYPOTHETICAL 47.4KD PROTEIN IN SHP1-SEC17 INTERGENIC REGION	536	42
730	gi7020242	Homo sapiens	cDNA FLJ20265 fis, clone COLF9334.	2813	99
730	gi14042159	Homo sapiens	cDNA FLJ14559 fis, clone NT2RM2001998.	2812	99
730	gi499005	Saccharomyce s cerevisiae	HRC830	128	32
731	gi7022375	Homo sapiens	cDNA FLJ10379 fis, clone NT2RM2002014.	3182	99
731	gi14010930	Homo sapiens	BAC clone RP11-576F1 from 2, complete sequence.	1868	100
731	gi1573555	Haemophilus influenzae Rd	transcription accessory protein (tex)	691	42
732	gi10434409	Homo sapiens	cDNA FLJ12737 fis, clone NT2RP2000337.	1001	99
733	gi7019597	Homo sapiens	clone PAC 270M7 chromosome 21 map 21q11.2, complete sequence.	5944	100
733	gi7407669	Homo sapiens	chromosome 21 PAC 30P13 map 21q11.2, complete sequence, containing gene for nuclear factor RIP140.	5944	100
733	gi7717256	Homo sapiens	chromosome 21 segment HS21C007.	5944	100
734	gi7021956	Homo sapiens	cDNA FLJ10103 fis, clone HEMBA1002495, weakly similar to LIGHT-MEDIATED DEVELOPMENT PROTEIN DET1.	1415	100
734	AAB64828	Homo sapiens	Human secreted protein sequence encoded by gene 12 SEQ ID NO:114.	869	99
734	gi4038594	Lycopersicon esculentum	tDET1 protein	413	37
735	gi6752405	Streptococcus pneumoniae	PspA	137	24
736	gi5080758	Homo sapiens	chromosome 19, BAC 331191 (CIT-B-471f3), complete sequence.	1486	55
736	gi456269	Mus musculus domesticus	zinc finger protein 30	1478	54

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
736	gi4567179	Homo sapiens	chromosome 19, BAC 37295 (CIT-B-21A4), complete sequence.	1281	62
737	gi7023220	Homo sapiens	cDNA FLJ10893 fis, clone NT2RP4002791.	4557	99
737	gi14042072	Homo sapiens	cDNA FLJ14507 fis, clone NT2RM1000399.	4439	97
737	gi7582296	Homo sapiens	BM-012	1807	99
738	gi11596985	Homo sapiens	chromosome 14 clone RP11-361H10 map 14q24.3, complete sequence.	1751	100
738	gi7020945	Homo sapiens	cDNA FLJ20689 fis, clone KAIA2890.	1738	99
738	gi6067151	Homo sapiens	chromosome 14 BAC 98L12, complete sequence.	1159	100
739	gi6941888	Homo sapiens	ubiquitin-specific processing protease (USP25) mRNA, complete cds.	5638	99
739	AAB31550	Homo sapiens	A human ubiquitin specific protease (USP).	5638	99
739	gi6693824	Homo sapiens	ubiquitin-specific protease (USP21) mRNA, complete cds.	4022	99
740	gi6693824	Homo sapiens	ubiquitin-specific protease (USP21) mRNA, complete cds.	5465	99
740	AAB31546	Homo sapiens	A human ubiquitin specific protease 25 (USP25).	5465	99
740	AAF24881_ aa1	Homo sapiens	DNA encoding a human ubiquitin specific protease 25 (USP25).	5465	99
741	gi7161175	Homo sapiens	mRNA for 19A24 protein (19A24 gene).	1726	100
741	gi13021810	Homo sapiens	NK cell receptor (CS1) mRNA, complete cds.	1349	100
741	AAB32373	Homo sapiens	Human secreted protein sequence encoded by gene 3 SEQ ID NO:59.	1349	100
742	gi7023747	Homo sapiens	cDNA FLJ11219 fis, clone PLACE1008122.	2553	100
742	gi7022222	Homo sapiens	cDNA FLJ10287 fis, clone HEMBB1001387.	880	97
742	AAG01392	Homo sapiens	Human secreted protein, SEQ ID NO: 5473.	569	99
743	gi7023747	Homo sapiens	cDNA FLJ11219 fis, clone PLACE1008122.	2442	97
743	gi7022222	Homo sapiens	cDNA FLJ10287 fis, clone HEMBB1001387.	769	89
743	AAG01392	Homo sapiens	Human secreted protein, SEQ ID NO: 5473.	569	99
744	gi6434857	Homo sapiens	pallid mRNA, complete cds.	872	100
744	gi13435969	Homo sapiens	, pallid (mouse) homolog, pallidin, clone MGC:4983, mRNA, complete cds.	872	100
744	gi6456870	Mus musculus	syntaxin 13-interacting protein pallid	754	87
745	gi6841480	Homo sapiens	HSPC129	2378	99
745	gi6841354	Homo sapiens	HSPC058	1825	99
745	gi7022613	Homo sapiens	cDNA FLJ10523 fis, clone NT2RP2000863.	1489	99
746	gi7023644	Homo sapiens	cDNA FLJ11155 fis, clone PLACE1006935.	1826	99
746	AAB18981	Homo sapiens	Amino acid sequence of a human transmembrane protein.	1000	99
746	gi13384531	Caenorhabditis elegans	similar to C. elegans protein T16H12.10	680	40
747	gi13544089	Homo sapiens	, clone IMAGE:4053618, mRNA, partial cds.	2749	99
747	gi6007859	Chlamydomon as reinhardtii	dynein heavy chain alpha	246	30

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
747	gi2065436	Schizosacchar omyces pombe	tealp	227	28
748	gi6650778	Homo sapiens	PRO1575	297	100
749	gi8926849	Homo sapiens	mRNA for Pex3p, complete cds.	1892	99
749	gi4092648	Homo sapiens	mRNA for PEX3 protein, partial.	1892	99
749	gi4218426	Homo sapiens	pex3 gene (joined CDS, promoter and exon 1).	1892	99
750	gi309209	Mus musculus	early B-cell factor	3064	99
750	gi6630994	Homo sapiens	early B-cell transcription factor (EBF) mRNA, partial cds.	3033	98
750	gi7687988	Gallus gallus	early B-cell factor	3023	97
751	gi10436636	Homo sapiens	cDNA FLJ14228 fis, clone NT2RP3004148.	3102	99
751	gi14278861	Homo sapiens	PHD zinc finger transcription factor mRNA, complete cds.	2127	100
751	gi12804495	Homo sapiens	, clone IMAGE:3356959, mRNA, partial cds.	1472	100
752	gi6594639	Homo sapiens	dynein intermediate chain DNAII (DNAII) mRNA, complete cds.	1773	100
752	gi6635422	Homo sapiens	dynein intermediate chain DNAII (DNAII) gene, exon 20 and complete cds.	1768	99
752	gi927637	Anthocidaris crassispina	dynein intermediate chain 2	961	61
753	gi5924385	Rattus norvegicus	ribosomal protein S271	412	100
753	gi12803647	Homo sapiens	, ribosomal protein S27 (metallopanstimulin 1), clone MGC:3659, mRNA, complete cds.	412	100
753	gi1373421	Homo sapiens	Human ribosomal protein S27 mRNA, complete cds.	412	100
754	gi1655432	Mus musculus	plexin 2	9646	96
754	gi6010215	Homo sapiens	mRNA for partial OCT/plexin-A2 protein.	6985	99
754	gi1665757	Mus musculus	plexin 1	6359	63
755	gi7770189	Homo sapiens	PRO2325	901	100
756	gi7022885	Homo sapiens	cDNA FLJ10697 fis, clone NT2RP3000527, weakly similar to ZINC FINGER PROTEIN 43.	3318	99
756	gi10434872	Homo sapiens	cDNA FLJ13043 fis, clone NT2RP3001338, weakly similar to ZINC FINGER PROTEIN 81.	957	43
756	gi38032	Homo sapiens	Human ZNF43 mRNA.	346	25
757	gi14042238	Homo sapiens	cDNA FLJ14604 fis, clone NT2RP1000363, moderately similar to R.norvegicus LL5 mRNA.	1107	93
757	AAB43723	Homo sapiens	Human cancer associated protein sequence SEQ ID NO:1168.	647	86
757	gi14044043	Homo sapiens	, clone IMAGE:4299555, mRNA, partial cds.	467	66
758	gi7106766	Homo sapieris	HSPC188	532	100
758	gi12804349	Homo sapiens	, clone MGC:4355, mRNA, complete cds.	529	99
758	gi1002516	Saccharomyce s cerevisiae	Hgh1p	115	27
759	gi6175593	Homo sapiens	transcription factor IIIC90 mRNA, complete cds.	4326	99
760	gi7023345	Homo sapiens	cDNA FLJ10970 fis, clone PLACE1000948.	647	99
760	AAG03409	Homo sapiens	Human secreted protein, SEQ ID NO:	239	100

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
	1		7490.		Auditity
761	gi5441541	Canis familiaris	Ribosomal protein	447	94
761	gi304526	Cricetulus griseus	ribosomal protein S17	447	94
761	gi10439453	Homo sapiens	cDNA: FLJ22917 fis, clone KAT06430.	447	94
762	gi6635353	Homo sapiens	RU1 (RU1) mRNA, complete cds.	4638	99
762	gi8100079	Mus musculus	polycomb-group proteins	4176	88
762	gi8100077	Rattus norvegicus	polycomb-group protein	4152	88
763	gi12804681	Homo sapiens	, \$100 calcium-binding protein, beta (neural), clone MGC:1323, mRNA, complete cds.	479	100
763	gi337730	Homo sapiens	Human \$100 protein beta-subunit gene, exon 3.	479	100
763	gi404769	Mus musculus	S100 beta protein	473	98
764	gi7106782	Homo sapiens	HSPC196	617	98
764	gi7106786	Homo sapiens	HSPC198	617	98
764	AAW74871	Homo sapiens	Human secreted protein encoded by gene 143 clone HBMDM46.	617	98
765	gi3851206	Homo sapiens	chromosome 19, cosmid F19847, complete sequence.	1282	100
765	gi13276629	Homo sapiens	mRNA; cDNA DKFZp761D221 (from clone DKFZp761D221); complete cds.	815	35
765	gi5701573	Caenorhabditis elegans	similar to S. pombe phosphoprotein (GB:X86179)	430	33
766	gi7020238	Homo sapiens	cDNA FLJ20262 fis, clone COLF7748.	1393	100
766	gi12653607	Homo sapiens	, clone IMAGE:3162218, mRNA, partial cds.	1019	98
766	AAY86358	Homo sapiens	Human gene 11-encoded protein fragment, SEQ ID NO:273.	996	95
767	gi2588619	Homo sapiens	BAC clone CTB-104F4 from 7q21-q22, complete sequence.	2037	100
767	gi1707507	Homo sapiens	H.sapiens mRNA for mitochondrial transcription termination factor.	2037	100
767	gi12654289	Homo sapiens	, transcription termination factor, mitochondrial, clone MGC:5000, mRNA, complete cds.	2033	99
768	gi1314373	Homo sapiens	Human aquaporin-5 (AQP5) gene, exon 4 and complete cds.	1336	100
768	gi664760	Rattus norvegicus	aquaporin-5	1245	91
768	gi4894460	Mus musculus	aquaporin 5	1235	91
769	gi13097624	Homo sapiens	, clone IMAGE:3608084, mRNA, partial cds.	1093	100
769	gi10438279	Homo sapiens	cDNA: FLJ22029 fis, clone HEP08661.	615	60
769	gi13325154	Homo sapiens	, clone IMAGE:3635709, mRNA, partial cds.	609	45
770	AAB48789	Homo sapiens	Human prostate cancer-predisposing protein, CA7 CG04.	2878	100
770	gi11321424	Mus musculus	Ral-A exchange factor RalGPS2	2073	96
770	gi7637906	Homo sapiens	Ral guanine nucleotide exchange factor RalGPS1A mRNA, complete cds.	1224	70
771	gi13623239	Homo sapiens	, Similar to SGC32445 protein, clone MGC:10610, mRNA, complete cds.	1080	99
771	gi7547035	Homo sapiens	SGC32445 protein (SGC32445) mRNA, complete cds.	687	100
771 ·	gi10434977	Homo sapiens	cDNA FLJ13110 fis, clone NT2RP3002549, moderately similar to	519	64

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
			HYPOTHETICAL 26.6 KD PROTEIN T19C3.4 IN CHROMOSOME III.		
772	gi13939858	Homo sapiens	RITA	2614	100
772	gi10048470	Homo sapiens	C2H2-like zinc finger protein (ZNF463) mRNA, complete cds.	2614	100
772	gi8575775	Homo sapiens	KRAB zinc finger protein (RITA) mRNA, complete cds.	2614	100
773	gi12654989	Homo sapiens	, clone MGC:5623, mRNA, complete cds.	2300	100
773	gi3329425	Homo sapiens	huntingtin interacting protein HYPE mRNA, partial cds.	963	100
773	gi429189	Haemophilus somnus	surface protein	152	41
774	gi14028017	Mesorhizobiu m loti	argininosuccinate lyase	199	26
774	gi2182606	Rhizobium sp. NGR234	Y4rH	179	29
775	gi3098311	Oryctolagus cuniculus	elongation factor 1 A2	2410	100
775	gi8886507	Homo sapiens	elongation factor 1 A-2 (EF1A-2) gene, complete cds.	2410	100
775	gi12653327	Homo sapiens	, eukaryotic translation elongation factor l alpha 2, clone MGC:8362, mRNA, complete cds.	2410	100
776	gi6624095	Homo sapiens	BAC clone RP11-294L11 from 2, complete sequence.	2515	97
776	AAY66674	Homo sapiens	Membrane-bound protein PRO1277.	2515	97
776	AAB87542	Homo sapiens	Human PRO1277.	2515	97
777	gi6049162	Homo sapiens	rhabdoid tumor deletion region protein 1 (RTDR1) mRNA, complete cds.	1732	100
777	gi14290442	Homo sapiens	, rhabdoid tumor deletion region protein 1, clone MGC:16968, mRNA, complete cds.	1732	100
778	AAB66071	Homo sapiens	Human INTERCEPT 296.	1787	99
778	AAB18992	Homo sapiens	Amino acid sequence of a human transmembrane protein.	880	58
778	AAB26325	Homo sapiens	Human CASB618 protein.	880	58
779	gi643656	Rattus norvegicus	synaptotagmin VII	1851	95
779	gi12667446	Rattus norvegicus	synaptotagmin VIIs	1851	95
779	gi6136786	Mus musculus	synaptotagmin VII	1842	95
780	gi7020988	Homo sapiens	cDNA FLJ20716 fis, clone HEP19742.	1048	100
780	gi4033606	Adiantum capillus- veneris	Extensin	131	38
780	gi169347	Phaseolus yulgaris	hydroxyproline-rich glycoprotein	130	38
781	gi7020477	Homo sapiens	cDNA FLJ20401 fis, clone KAT00901.	1644	96
781	gi7022002	Homo sapiens	cDNA FLJ10135 fis, clone HEMBA1003117.	590	40
781	gi7022284	Homo sapiens	cDNA FLJ10324 fis, clone NT2RM2000567.	590	40
782	gi6808186	Homo sapiens	mRNA; cDNA DKFZp434D0218 (from clone DKFZp434D0218); partial cds.	1322	99
783	gi505544	Homo sapiens	H.sapiens mRNA for Zinc-finger protein (ZNFpT1).	1211	99
783	AAY58627	Homo sapiens	Protein regulating gene expression PRGE-20.	688	50
783	gi9187356	Homo sapiens	mRNA full length insert cDNA clone	687	50

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
			EUROIMAGE 2107395.		
784	gi8896094	Homo sapiens	SH3-containing protein SH3GLB2 mRNA, complete cds.	1975	97
784	gi4929591	Homo sapiens	CGI-61 protein mRNA, complete cds.	706	69
784	gi8896092	Homo sapiens	SH3-containing protein SH3GLB1 mRNA, complete cds.	706	69
785	gi7770175	Homo sapiens	PRO2249	1827	99
785	gi11527602	Homo sapiens	mRNA for MCM10 homolog, complete cds.	1827	99
785	gi12053187	Homo sapiens	mRNA; cDNA DKFZp434H152 (from clone DKFZp434H152); complete cds.	1682	99
786 .·	gi7023364	Homo sapiens	cDNA FLJ10982 fis, clone PLACE1001692, moderately similar to S- ACYL FATTY ACID SYNTHASE THIOESTERASE, MEDIUM CHAIN (EC 3.1.2.14).	1413	99
786	gi7023563	Homo sapiens	cDNA FLJ11106 fis, clone PLACE1005763, moderately similar to S- ACYL FATTY ACID SYNTHASE THIOESTERASE, MEDIUM CHAIN (EC 3.1.2.14).	1099	98
786	gi205326	Rattus norvegicus	S-acyl fatty acid sunthetase thio ester hydrolase, medium chain	-807	55
787	gi2599502	Homo sapiens	protocadherin 68 (PCH68) mRNA, complete cds.	327	43
787	AAY24913	Homo sapiens	Human ontherin.	327	43
787	AAY94991	Homo sapiens	Human secreted protein vc35_1, SEQ ID NO:22.	296	28
788	gi7023688	Homo sapiens	cDNA FLJ11183 fis, clone PLACE1007488, weakly similar to PUTATTVE RHO/RAC GUANINE NUCLEOTIDE EXCHANGE FACTOR.	2260	100
788	gi3342246	Rattus norvegicus	actin-filament binding protein Frabin	725	32
788	gi595425	Homo sapiens	Human faciogenital dysplasia (FGD1) mRNA, complete cds.	759	32
789	gi6554165	Homo sapiens	receptor protein tyrosine phosphatase (RPTP-rho) mRNA, alternatively spliced, complete cds.	7734	99
789	gi13378306	Mus musculus	brain RPTPmam4 isoform I	7499	97
789	gi32456	Homo sapiens	H.sapiens hR-PTPu gene for protein tyrosine phosphatase.	4995	64
790	gi7020479	Homo sapiens	cDNA FLJ20402 fis, clone KAT00919.	2024	99
790	gi7770205	Homo sapiens	PRO2521	1957	97
790	gi10241843	Mus musculus	gasdermin	282	29
791	gi5262472	Homo sapiens	mRNA; cDNA DKFZp564J102 (from clone DKFZp564J102); partial cds.	1602	100
792	gi10436457	Homo sapiens	cDNA FLJ14084 fis, clone HEMBB1002383.	830	100
792	AAY94940	Homo sapiens	Human secreted protein clone yi62_1 protein sequence SEQ ID NO:86.	830	100
792	AAY57922	Homo sapiens	Human transmembrane protein HTMPN-46.	830	100
793	gi7328061	Homo sapiens	mRNA; cDNA DKFZp761I2312 (from clone DKFZp761I2312); partial cds.	2723	100
793	gi14039825	Mus musculus	gamma-1 syntrophin	2579	93
793	gi8247279	Homo sapiens	mRNA for syntrophin 4.	2271	97
794	gi6164674	Homo sapiens	heterogeneous nuclear ribonucleoprotein, alternate transcript (RALY) mRNA,	730	66

SEQ ID	Accession No.	Species	Description	Score	% Identity
N.O.	110.	 	complete cds.	 	Adentity
794	gi14250048	Homo sapiens	, heterogeneous nuclear ribonucleoprotein C (C1/C2), clone MGC:14574, mRNA, complete cds.	705	53
794	gi13937888	Homo sapiens	, Similar to heterogeneous nuclear ribonucleoprotein C, clone MGC:12469, mRNA, complete cds.	704	53
795	gi12653905	Homo sapiens	, Similar to Max dimerization protein 3, clone MGC:2383, mRNA, complete cds.	1045	100
795	AAY93137	Homo sapiens	Human Myx protein.	1023	98
795	AAB35713	Homo sapiens	Human Mad3 protein sequence.	1010	97
796	gi7020704	Homo sapiens	cDNA FLJ20533 fis, clone KAT10931.	585	98
797	gi7106878	Homo sapiens	HSPC244	398	98
797	AAY07855	Homo sapiens	Human secreted protein fragment encoded from gene 4.	398	98
797	gi13274582	Mus musculus	thymus atrophy-related protein	383	95
798	gi8886483	Gallus gallus	EURL	1178	74
798	gi10435877	Homo sapiens	cDNA FLJ13763 fis, clone PLACE4000089.	873	98
798	AAG01108	Homo sapiens	Human secreted protein, SEQ ID NO: 5189.	561	100
799	AAY33297	Homo sapiens	Human membrane spanning protein MSP-4.	.781	100
799	AAB61149	Homo sapiens	Human NOV18 protein.	781	100
799	AAB61150	Homo sapiens	Human NOV19 protein.	781	100
800	gi8099348	Homo sapiens	zinc finger protein (ZFP) mRNA, complete cds.	4066	99
800	gi2293535	Homo sapiens	zinc finger protein (ZnF20) mRNA, complete cds.	1863	49
800	gi11527849	Mus musculus	zinc finger protein SKAT2	1323	58
801	gi7023523	Homo sapiens	cDNA FLJ11082 fis, clone PLACE1005206.	2693	99
801	gi9558010	Leishmania major	possible cDNA flj11082 fis, clone place1005206	134	26
802	gi6841558	Homo sapiens	HSPC168	1502	100
802	gi6453346	Homo sapiens	Novel human gene on chromosome 20.	1502	100
802	gi13542748	Mus musculus	RIKEN cDNA 3230401D17 gene	1314	86
803	gi7020468	Homo sapiens	cDNA FLJ20396 fis, clone KAT00561.	931	100
803	AAB18980	Homo sapiens	Amino acid sequence of a human transmembrane protein.	931	100
803	AAY91632	Homo sapiens	Human secreted protein sequence encoded by gene 25 SEQ ID NO:305.	914	98
804	gi6650345	Homo sapiens	alpha-catenin-like protein VR22 mRNA, complete cds.	4478	99
804	gi222788	Gallus gallus	alpha N-catenin	2765	60
804	AAR58778	Homo sapiens	Neural alpha-catenin protein.	2765	60
805	gi10434911	Homo sapiens	cDNA FLJ13068 fis, clone NT2RP3001739, weakly similar to HYPOTHETICAL 72.5 KD PROTEIN C2F7.10 IN CHROMOSOME I.	587	38
805	gi5912258	Homo sapiens	mRNA; cDNA DKFZp586K0524 (from clone DKFZp586K0524); partial cds.	190	41
805	gi7022673	Homo sapiens	cDNA FLJ10562 fis, clone NT2RP2002701.	154	44
806	gi10435877	Homo sapiens	cDNA FLJ13763 fis, clone PLACE4000089.	876	99
806	gi8886483	Gallus gallus	EURL	868	72
806	AAG01108	Homo sapiens	Human secreted protein, SEQ ID NO:	561	100
	I	1	5189.		

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
807	gi4521254	Mus musculus	cornichon-like protein	867	100
807	AAB60464	Homo sapiens	Human cell cycle and proliferation protein CCYPR-12, SEQ ID NO:12.	729	81
807	AAY76218	Homo sapiens	Human secreted protein encoded by gene 95.	716	81
808	gi7407144	Homo sapiens	protocadherin Fat 2 (FAT2) mRNA, complete cds.	22667	99
808	gi3449286	Rattus norvegicus	MEGF1	18806	81
808	gi6688786	Mus musculus	mouse fat 1 cadherin	8928	47
809	gi7407144	Homo sapiens	protocadherin Fat 2 (FAT2) mRNA, complete cds.	19770	99
809	gi3449286	Rattus norvegicus	MEGF1 ·	16567	82
809	gi6688786-	Mus musculus	mouse fat 1 cadherin	8928	47
810	gi7020201	Homo sapiens	cDNA FLJ20241 fis, clone COLF6335.	2420	100
810	gi10435321	Homo sapiens	cDNA FLJ13337 fis, clone OVARC1001880.	1279	99
810	gi7020600	Homo sapiens	cDNA FLJ20475 fis, clone KAT07206.	634	60
811	gi6483290	Homo sapiens	CDH7 mRNA for cadherin-7, complete cds.	4032	100
811	gi10803408	Homo sapiens	mRNA for cadherin-7 (CDH7 gene).	3965	98
811	gi868001	Gallus gallus	chicken cadherin-7	3830	93
812	gi13276621	Homo sapiens	mRNA; cDNA DKFZp761G1913 (from clone DKFZp761G1913).	1204	97
812	gi8977983	Mus musculus	neuronal interacting factor X 1 (NIX1)	699	78
812	gi10437116	Homo sapiens	cDNA: FLJ21097 fis, clone CAS03931.	297	42
814	gi13279269	Homo sapiens	, clone IMAGE:3631943, mRNA, partial cds.	1480	100
814	gi6808028	Homo sapiens	mRNA; cDNA DKFZp761C029 (from clone DKFZp761C029); partial cds.	857	100
814	AAW88657	Homo sapiens	Secreted protein encoded by gene 124 clone HPMCJ92.	436	94
815	gi7959853	Homo sapiens	PRO1966	281	100
816	gi7259234	Mus musculus	contains transmembrane (TM) region	718	65
816	AAY94954	Homo sapiens	Human secreted protein clone iw66_1 protein sequence SEQ ID NO:114.	679	58
816	AAB62810	Homo sapiens	Human nervous system associated protein NSPRT3 amino acid sequence.	678	61
817	gi5921144	Schizosacchar omyces pombe	mip1	1489	48
817	gi458938	Saccharomyce s cerevisiae	Yhr186cp	469	30
817	gi9366720	Trypanosoma brucei	possible t16o11.22 protein.	277	45
819	gi7020799	Homo sapiens	cDNA FLJ20590 fis, clone KAT09052.	727	100
820	gi7020555	Homo sapiens	cDNA FLJ20449 fis, clone KAT05575.	1857	99
820	AAY79269	Homo sapiens	Human testis-specific transcription factor PHELIX.	1696	99
821	gi6482350	Homo sapiens	CAC-1 mRNA, partial cds.	1136	100
821	gi13937595	Homo sapiens	, Similar to RIKEN cDNA 1810017F10 gene, clone MGC:2583, mRNA, complete cds.	560	94
821	AAY25770	Homo sapiens	Human secreted protein encoded from gene 60.	560	94
822	gi10434608	Homo sapiens	cDNA FLJ12871 fis, clone NT2RP2003751.	2023	100
822	gi6093227	Homo sapiens	mRNA; cDNA DKFZp434I0850 (from clone DKFZp434I0850); partial cds.	1607	100

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
822	gi6453452	Homo sapiens	mRNA; cDNA DKFZp434L0850 (from clone DKFZp434L0850).	1607	100
823	AAY13402	Homo sapiens	Amino acid sequence of protein PRO310.	1079	63
823	AAB18988	Homo sapiens	Amino acid sequence of a human transmembrane protein.	1079	63
823	AAB80270	Homo sapiens	Human PRO310 protein.	1079	63
824	gi13938181	Homo sapiens	, clone IMAGE:2905978, mRNA, partial cds.	2722	99
824	gi6453540	Homo sapiens	mRNA; cDNA DKFZp434D0428 (from clone DKFZp434D0428); partial cds.	2455	99
824	gi10440436	Homo sapiens	mRNA for FLJ00053 protein, partial cds.	807	100
825	gi7022318	Homo sapiens	cDNA FLJ10346 fis, clone NT2RM2001004.	1475	100
826	gi7110152	Mus musculus	selenocysteine lyase SCLY	1219	80
826	gi7022600	Homo sapiens	cDNA FLJ10515 fis, clone NT2RP2000764, weakly similar to NIFS PROTEIN.	592	98
826	gi9887215	Methanosarcin a thermophila	cysteine desulfurase NifS	315	43
827	gi7022560	Homo sapiens	cDNA FLJ10491 fis, clone NT2RP2000239.	1266	100
827	gi7022033	Homo sapiens	cDNA FLJ10156 fis, clone HEMBA1003447.	1161	97
828 ⁻	gi8247250	Homo sapiens	mRNA for neutral sphingomyelinase II (nSMase2 gene).	3489	100
828	AAB70772	Homo sapiens	Human neutral cerebral sphingomyelinase protein.	3489	100
828	gi8247281	Mus musculus	neutral sphingomyelinase II	3187	91
829	gi7020945	Homo sapiens	cDNA FLJ20689 fis, clone KAIA2890.	2459	100
829	gi11596985	Homo sapiens	chromosome 14 clone RP11-361H10 map 14q24.3, complete sequence.	1819	97
829	gi6067151	Homo sapiens	chromosome 14 BAC 98L12, complete sequence.	1153	99
830	gi10039443	Homo sapiens	NEDL1 mRNA for NEDD4-like ubiquitin ligase 1, complete cds.	4335	56
830	AAW93167	Homo sapiens	Human ZGGBP1 protein.	992	47
830	gi1374782	Mus musculus	possible ubiquitin protein ligase	1062	50
831	gi7021974	Homo sapiens	cDNA FLJ10115 fis, clone HEMBA1002777.	1882	99
831	gi7021027	Homo sapiens	cDNA FLJ20739 fis, clone HEP07341.	1252	98
831	gi5002381	Takifugu rubripes	BAW	776	72
832	gi7022523	Homo sapiens	cDNA FLJ10469 fis, clone NT2RP2000008, weakly similar to ZINC FINGER PROTEIN 84.	3772	99
832	gi1020145	Homo sapiens	Human DNA binding protein (HPF2) mRNA, complete cds.	1714	48
832	gi7243633	Homo sapiens	RB-associated KRAB repressor (RBAK) mRNA, complete cds.	1653	46
833	gi6433864	Homo sapiens	CLDN12 gene for claudin-12.	1295	100
833	gi12053057	Homo sapiens	mRNA; cDNA DKFZp434I1816 (from clone DKFZp434I1816); complete cds.	1295	100
833	gi9799020	Mus musculus	claudin-12	1125	91
834	gi12053151	Homo sapiens	mRNA; cDNA DKFZp434G0326 (from clone DKFZp434G0326); complete cds.	5605	99
834	gi7020102	Homo sapiens	cDNA FLJ20176 fis, clone COL09928.	1268	88
834	gi7020102 gi7023725	Homo sapiens	cDNA FLJ11205 fis, clone PLACE1007843.	719	100
835	gi7020789	Homo sapiens	cDNA FLJ20583 fis, clone KAT09685.	2153	99

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
835	AAG02503	Homo sapiens	Human secreted protein, SEQ ID NO: 6584.	423	98
835	gi14289183	Homo sapiens	chac mRNA for chorein, complete cds.	193	24
836	gi7022600	Homo sapiens	cDNA FLJ10515 fis, clone NT2RP2000764, weakly similar to NIFS PROTEIN.	1301	100
836	gi7110152	Mus musculus	selenocysteine lyase SCLY	1107	83
836	gi13592392	Caenorhabditis elegans	Contains similarity to Pfam domain: PF00266 (aminotran_5), Score=51.6, E-value=5.7e-12, N=1	468	44
837	gi7274380	Homo sapiens	group III secreted phospholipase A2 mRNA, complete cds.	2813	99
837	gi4314431	Homo sapiens	PAC clone RP3-412A9 from 22, complete sequence.	596	99
837	gi5627	Apis mellifera	phospholipase A-2	243	41
	gi8331760	Homo sapiens	X28 region near ALD locus containing dual specificity phosphatase 9 (DUSP9), ribosomal protein L18a (RPL18a), Ca2+/Calmodulin-dependent protein kinase I (CAMKI), creatine transporter (CRTR), CDM protein (CDM), adrenoleukodystrophy protein (ALD), plexin-related protein (PLXB3), musclespecific serine kinase (MSSK), NADisocitrate dehydrogenase (IDH), translocon-associated protein delta (TRAP), and LUI protein (LUI) genes, complete cds; and CCp pseudogene, complete sequence.	3242	100
838	gi6651019	Mus musculus	semaphorin cytoplasmic domain- associated protein 3A	1583	50
838	gi6651021	Mus musculus	semaphorin cytoplasmic domain- associated protein 3B	1583	50
839	gi7023290	Homo sapiens	cDNA FLJ10932 fis, clone OVARC1000588.	718	100
840	gi6094681	Homo sapiens	PAC clone RP5-1049N15 from 7q31.2-7q32, complete sequence.	4804	100
840	gi7264724	Homo sapiens	alpha-aminoadipate semialdehyde synthase mRNA, complete cds.	4804	100
840	gi4938304	Homo sapiens	mRNA for lysine-ketoglutarate reductase/saccharopine dehydrogenase, partial CDS.	4799	99
841	AAY66700	Homo sapiens	Membrane-bound protein PRO1137.	1164	95
841	AAB65223	Homo sapiens	Human PRO1137 (UNQ575) protein sequence SEQ ID NO:250.	1164	95
841	AAY50917	Homo sapiens	Human fetal brain cDNA clone vc4_1 derived protein.	1023	100
842	AAW56477	Homo sapiens	Amino acid sequence of human bone morphogenetic protein-16 (BMP-16).	1183	100
842	AAY03849	Homo sapiens	Human nodal protein.	1183	100
842	gi296605	Mus musculus	nodal	986	84
843	gi7020399	Homo sapiens	cDNA FLJ20356 fis, clone HEP15821.	5470	100
843	gi10435659	Homo sapiens	cDNA FLJ13605 fis, clone PLACE1010562.	224	44
844	gi4886471	Homo sapiens	mRNA; cDNA DKFZp586N0819 (from clone DKFZp586N0819).	531	100
845	gi3288470	Homo sapiens	surf5c mRNA, clone 10.9.	728	100
845	gi3288452	Homo sapiens	Surf-5 and Surf-6 genes.	334	94
845	gi3288468	Homo sapiens	surf5b mRNA, clone L5.	334	94
846	gi14149050	Drosophila	turtle protein, isoform 4	1037	32

SEQ ID	Accession No.	Species	Description	Score	% Identity
		melanogaster			
846	gi14149048	Drosophila melanogaster	turtle protein, isoform 3	1037	32
846	gi14149046	Drosophila melanogaster	turtle protein, isoform 2	939	34
847	gi7021049	Homo sapiens	cDNA FLJ20753 fis, clone HEP02714.	2930	99
847	gi9886896	Human herpesvirus 8	Orf73	175	20
847	gi11037008	Human herpesvirus 8	latent nuclear antigen	172	20
848	gi12597293	Homo sapiens	acidic mammalian chitinase precursor, mRNA, complete cds.	2018	100
848	gi6467177	Homo sapiens	TSA1902-L mRNA for novel member of chitinase family, complete cds.	2010	99
848	gi6467179	Homo sapiens	TSA1902-S mRNA for novel member of chitinase family, complete cds.	1725	99
849	gi32391	Homo sapiens	Human HOX4C mRNA for a homeobox protein.	1802	98
849	gi51416	Mus musculus	Hox-4.4	1591	88
849	gi4322104	Danio rerio	homeobox protein	425	82
850	gi1359443	Homo sapiens	Human gene for hepatitis C-associated microtubular aggregate protein p44, exon 9 and complete cds.	2299	99
850	AAY05371	Homo sapiens	Human HCMV inducible gene protein, SEQ ID NO 10.	2299	99
850	gi218576	Pan troglodytes	p44	2242	97
851	gi575494	Homo sapiens	MHC class II lymphocyte antigen beta- chain (HLA-DPB1a) mRNA, complete cds.	437	72
851	gi188479	Homo sapiens	Human MHC class II lymphocyte antigen (HLA-DP) beta chain mRNA, complete cds.	437	72
851	gi14044082	Homo sapiens	, Similar to major histocompatibility complex, class II, DP beta 1, clone MGC:14112, mRNA, complete cds.	429	70
852	gi181547	Homo sapiens	defensin 6 mRNA, complete cds.	318	90
852	AAR44819	Homo sapiens	Sequence of the gastrointestinal defensin (GID) peptide calledhuman defensin 6.	318	90
852	gi1200182	Homo sapiens	Human defensin 6 (HD-6) gene, complete cds.	314	89
853	gi13396914	Homo sapiens	The gene of C2GnT3	2389	100
853	gi7527464	Homo sapiens	core 2 beta-1,6-N- acetylglucosaminyltransferase 3 (C2GnT3) mRNA, complete cds.	2389	100
853	AAU00037	Homo sapiens	Human C2GnT3.	2389	100
855	gi7959772	Homo sapiens	PRO1483	252	100
856	gi5911169	Homo sapiens	transmembrane mucin 12 (MUC12) mRNA, partial cds.	2914	99
856	AAY59290	Homo sapiens	Human MUC12 polypeptide.	2914	99
856	gi2589172	Rattus norvegicus	mucin Muc3	595	36
857	AAE00508	Homo sapiens	Human lipase protein, MLip-1.	1456	100
857	gi56600	Rattus norvegicus	triacylglycerol lipase	776	58
857	gi3108175	Mus musculus	pancreatic lipase related protein 1	772	57
858	AAY94954	Homo sapiens	Human secreted protein clone iw66_1 protein sequence SEQ ID NO:114.	1112	100
858	gi10434269	Homo sapiens	cDNA FLJ12650 fis, clone	872	100

SEQ ID NO:	Accession No.	Species	Description	Score	. % Identity
NO:	140.		NT2RM4002054.	 	lucitity
858	gi7259234	Mus musculus	contains transmembrane (TM) region	660	60
859	gi7021851	Homo sapiens	cDNA FLJ10035 fis, clone HEMBA1000919.	1589	100
859	gi10440420	Homo sapiens	mRNA for FLJ00045 protein, partial cds.	654	89
859	AAY99671	Homo sapiens	Human GTPase associated protein-22.	654	89
860	gi7022523	Homo sapiens	cDNA FLJ10469 fis, clone NT2RP2000008, weakly similar to ZINC FINGER PROTEIN 84.	3573	99
860	gi1020145	Homo sapiens	Human DNA binding protein (HPF2) mRNA, complete cds.	1604	48
860	gi12584159	Homo sapiens	zinc finger protein 268 (ZNF268) mRNA, complete cds.	1542	48
861	gi6539434	Homo sapiens	SPR1 mRNA, complete cds.	808	100
861	gi6523547	Volvox carteri f. nagariensis	hydroxyproline-rich glycoprotein DZ- HRGP	185	39
861	gi904359	Beta vulgaris	chitinase l	185	41
862	gi7021924	Homo sapiens	cDNA FLJ10081 fis, clone HEMBA1002018.	2742	100
862	gi10435862	Homo sapiens	cDNA FLJ13751 fis, clone PLACE3000339, weakly similar to GLUCOAMYLASE S1/S2 PRECURSOR (EC 3.2.1.3).	2687	99
862	gi11275988	Homo sapiens	testis development protein PRTD mRNA, complete cds.	2454	99
863	gi7019913	Homo sapiens	cDNA FLJ20060 fis, clone COL01358.	1830	100
863	gi10434817	Homo sapiens	cDNA FLJ13006 fis, clone NT2RP3000449.	1823	.99
863	gi10434659	Homo sapiens	cDNA FLJ12902 fis, clone NT2RP2004347.	1724	99
864	gi7329718	Homo sapiens	Novel human gene mapping to chomosome 1.	11682	99
864	gi7022765	Homo sapiens	cDNA FLJ10619 fis, clone NT2RP2005472.	3153	99
864	gi14388939	Homo sapiens	chorea-acanthocytosis (CHAC) mRNA, complete cds.	462	30
865	gi28971	Homo sapiens	H.sapiens mRNA for autoantigen NOR- 90.	3813	100
865	gi509241	Homo sapiens	Human mRNA for upstream binding factor (hUBF).	2661	78
865	AAB44430	Homo sapiens	Human lung tumour-specific antigen encoded by cDNA	2649	78
866	gi13445482	Homo sapiens	HP43.8KD mRNA, complete cds.	282	47
866	gi10434108	Homo sapiens	cDNA FLJ12552 fis, clone NT2RM4000712, moderately similar to Homo sapiens ubiquitin hydrolyzing enzyme I (UBH1) mRNA.	219	36
866	gi10436670	Homo sapiens	cDNA FLJ14256 fis, clone PLACE1000007, weakly similar to PROBABLE UBIQUITIN CARBOXYL- TERMINAL HYDROLASE R10E11.3 (EC 3.1.2.15).	219	36
867	AAB73229	Homo sapiens	Human phosphatase MTMR7 h	743	57
867	gi5901814	Drosophila melanogaster	BcDNA.GH04637	503	48
867	gi7020021	Homo sapiens	cDNA FLJ20126 fis, clone COL06160.	697	73
868	gi7959801	Homo sapiens	PRO0800	392	100
869	gi12654971	Homo sapiens	, calcium-regulated heat-stable protein (24kD), clone MGC:5586, mRNA,	417	97

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
			complete cds.		
869	gi13097198	Homo sapiens	, calcium-regulated heat-stable protein (24kD), clone MGC:5235, mRNA, complete cds.	417	97
869	AAW61023	Homo sapiens	Human RNA binding protein.	417	97
870	gi6650832	Homo sapiens	PRO2086	243	100
871	gi2217942	Rattus norvegicus	glycoprotein specific UDP- glucuronyltransferase	1802	97
871	gi8051678	Homo sapiens	hu-GlcAT-P mRNA for glucuronyltransferase, complete cds.	1757	99
871	gi4519214	Rattus norvegicus	UDP-glucuronyltransferase-S	760	50
872	gi14286288	Homo sapiens	, Similar to RIKEN cDNA 2010004P11 gene, clone MGC:2734, mRNA, complete cds.	715	100
872	gi13529665	Mus musculus	RIKEN cDNA 2010004P11 gene	706	98
872	gi2565364	Musca domeștica	Sex-lethal protein	134	33
873	gi190406	Homo sapiens	Human profilaggrin gene exons 1-3, 5' end.	6301	99
873	gi190396	Homo sapiens	Human profilaggrin gene, 3' end.	5133	99
873	gi190404	Homo sapiens	Human profilaggrin mRNA, 3' end.	3696	89
874	gi791002	Homo sapiens	ARSD gene, complete CDS.	1761	99
874	gi6651286	Homo sapiens	arylsulfatase D beta (ARSD) mRNA, complete cds.	1756	99
874	gi791004	Homo sapiens	ARSE gene, complete CDS.	947	58
875	gi13097675	Homo sapiens	, Similar to uncharacterized hypothalamus protein HCDASE, clone MGC:1171, mRNA, complete cds.	612	96
875	AAY87599	Homo sapiens	Human fatty acid beta-oxidation enzyme HUFA-2.	612	96
875	AAG03352	Homo sapiens	Human secreted protein, SEQ ID NO: 7433.	591	100
876	gi6180180	Homo sapiens	transcription factor IGHM enhancer 3, JM11 protein, JM4 protein, JM5 protein, T54 protein, JM10 protein, A4 differentiation-dependent protein, triple LIM domain protein 6, and synaptophysin genes, complete cds; and L-type calcium channel alpha-1 subunit gene, partial cds, complete sequence.	908	100
876	gi3114826	Homo sapiens	mRNA for JM4 protein, complete CDS (clone IMAGE 546750 and LLNLc110F1857Q7 (RZPD Berlin)).	908	100
876	gi7673612	Mus musculus	DXImx39e protein	831	91
877	gi13543663	Homo sapiens	, ubiquitin-conjugating enzyme E2D 1 (homologous to yeast UBC4/5), clone MGC:14673, mRNA, complete cds.	805	100
877	gi460810	Homo sapiens	H.sapiens UBCH5 mRNA for ubiquitin conjugating enzyme.	805	100
877	gi4868140	Homo sapiens	ubiquitin-conjugating enzyme HBUCE1 mRNA, complete cds.	747	91
878	gi7020915	Homo sapiens	cDNA FLJ20666 fis, clone KAIA608.	1288	100
878	gi3169096	Schizosacchar omyces pombe	possible pre-mRNA processing by similarity to yeast prp39	279	33
878	gi10177721	Arabidopsis thaliana	gene_id:MPL12.20~	146	22
879	gi7020681	Homo sapiens	cDNA FLJ20519 fis, clone KAT10365.	891	100
879	AAY87267	Homo sapiens	Human signal peptide containing protein	824	95

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
			HSPP-44 SEQ ID NO:44.		
879	AAB65245	Homo sapiens	Human PRO1104 (UNQ547) protein sequence SEQ ID NO:297.	824	95
880	gi6560622	Homo sapiens	PRO0611	501	100
881	AAB57079	Homo sapiens	Human prostate cancer antigen protein sequence SEQ ID NO:1657.	668	100
881	AAY99372	Homo sapiens	Human PRO1430 (UNQ736) amino acid sequence SEQ ID NO:116.	668	100
881	AAB88356	Homo sapiens	Human membrane or secretory protein clone PSEC0082.	661	99
882	gi1381181	Oryctolagus cuniculus	ubiquitin-conjugating enzyme E2-32k	663	100
882	gi13436071	Homo sapiens	, clone MGC:10481, mRNA, complete cds.	663	100
882	gi7020506	Homo sapiens	cDNA FLJ20419 fis, clone KAT02435.	658	99
883	gi1381181	Oryctolagus cuniculus	ubiquitin-conjugating enzyme E2-32k	1265	99
883	gi13436071	Homo sapiens	, clone MGC:10481, mRNA, complete cds.	1265	.99
883	gi7020506	Homo sapiens	cDNA FLJ20419 fis, clone KAT02435.	1256	98
884	gi1381181	Oryctolagus cuniculus	ubiquitin-conjugating enzyme E2-32k	383	97
884	gi13436071	Homo sapiens	, clone MGC:10481, mRNA, complete cds.	383	97
884	gi7020506	Homo sapiens	cDNA FLJ20419 fis, clone KAT02435.	383	97
885	gi14424536	Homo sapiens	, Similar to septin 6, clone MGC:16619, mRNA, complete cds.	2183	99
885	gi5689158	Mus musculus	Septin6	2114	95
885	gi7023141	Homo sapiens	cDNA FLJ10849 fis, clone NT2RP4001414, highly similar to SEPTIN 2 HOMOLOG.	1840	82
886	gi14424536	Homo sapiens	, Similar to septin 6, clone MGC:16619, mRNA, complete cds.	1213	63
886	gi5689158	Mus musculus	Septin6	1162	62
886	gi7023141	Homo sapiens	cDNA FLJ10849 fis, clone NT2RP4001414, highly similar to SEPTIN 2 HOMOLOG.	995	51
887	gi4309951	Homo sapiens	BAC clone RP11-121A8 from 7p14-p13, complete sequence.	684	100
887	AAG00417	Homo sapiens	Human secreted protein, SEQ ID NO: 4498.	684	100
887	gi339159	Homo sapiens	Human T-cell receptor germline gamma- chain gene V-region (V3; subgroup I).	392	73
888	gi2570015	Homo sapiens	H.sapiens PAX7 gene, exon 1 (and joined CDS).	2756	100
888	gi2570021	Homo sapiens	H.sapiens mRNA for paired box containing transcription factor, PAX7.	2756	100
888	gi2570014	Homo sapiens	H.sapiens PAX7 gene, exon 1 (and joined CDS).	2735	99

TABLE 3

TABLE 3			
SEQ ID NO:	Accession No.	Description	Results*
445	BL00434	HSF-type DNA-binding domain proteins.	BL00434C 23.85 7.111e-09 1089-1129
446	PD00066	PROTEIN ZINC-FINGER	PD00066 13.92 1.000e-13 216-229
		METAL-BINDI.	PD00066 13.92 2.286e-12 244-257
	•		PD00066 13.92 4.522e-11 299-312
	ļ		PD00066 13.92 6.538e-10 157-170
			PD00066 13.92 7.923e-10 327-340
453	PR00037	LACR BACTERIAL REGULATORY PROTEIN HTH SIGNATURE	PR00037A 12.66 6.786e-09 34-49
465	PR00320	G-PROTEIN BETA WD-40	PR00320C 13.01 6.100e-09 197-212
		REPEAT SIGNATURE	PR00320C 13.01 6.400e-09 393-408
	1		PR00320A 16.74 8.683e-09 197-212
			PR00320B 12.19 9.775e-09 299-314
466	DM00215	PROLINE-RICH PROTEIN 3.	DM00215 19.43 5.881e-09 14-47
470	BL00175	Phosphoglycerate mutase	BL00175D 27.67 8.500e-40 175-227
		family phosphohistidine	BL00175C 23.75 5.000e-25 90-122
	l	proteins.	BL00175A 15.42 8.333e-20 17-37
	ļ		BL00175B 12.60 1.000e-12 66-79
472	BL00315	Dehydrins proteins.	BL00315A 9.35 8.119e-09 105-133
473	BL00518	Zinc finger, C3HC4 type	BL00518 12.23 4.000e-11 44-53
		(RING finger), proteins.	
475	PD02448	TRANSCRIPTION	PD02448A 9.37 4.293e-09 171-210
		PROTEIN DNA-BINDIN.	PROCESS 10.04.0.500- 10.121.141
477	PR00625	DNAJ PROTEIN FAMILY	PR00625A 12.84 8.500e-19 121-141
		SIGNATURE	PR00625B 13.48 3.204e-15 151-172 PD02102A 16.74 5.853e-10 26-70
478	PD02102	SUBUNIT E V-ATPASE	PD02102A 10.74 5.8558-10 20-70
		VACUOLAR ATP	
450	BL00018	SYNTHASE HYDROL. EF-hand calcium-binding	BL00018 7.41 4.706e-11 49-62
479	BLUUUIS	domain proteins.	BB00016 7.41 4.7000 17 17 02
480	PR00501	KELCH REPEAT	PR00501A 8.25 9.182e-09 544-558
460	FROODU	SIGNATURE	110000111011101111111111111111111111111
483	PR00878	CHOLINESTERASE	PR00878F 5.37 5.179e-12 500-513
703	11100070	SIGNATURE	
484	BL00378	Hexokinases proteins.	BL00378C 16.14 1.000e-40 207-251
1.0.		F	BL00378E 22.92 1.000e-40 725-771
			BL00378C 16.14 3.520e-40 655-699
	1		BL00378E 22.92 3.382e-36 277-323
		1	BL00378B 14.23 5.333e-35 509-546
			BL00378B 14.23 8.953e-28 61-98
			BL00378A 19.01 1.346e-22 22-50
		1	BL00378F 8.27 2.688e-17 893-908
	1		BL00378D 10.94 6.294e-17 703-715
			BL00378D 10.94 5.500e-16 255-267
			BL00378F 8.27 9.609e-13 445-460
			BL00378A 19.01 3.017e-12 470-498
485	BL00028	Zinc finger, C2H2 type,	BL00028 16.07 2.688e-15 352-369
		domain proteins.	BL00028 16.07 4.375e-15 324-341
	ł		BL00028 16.07 4.176e-14 604-621
1	1	1	BL00028 16.07 8.412e-14 380-397
	1	1	BL00028 16.07 9.471e-14 576-593
1		1	BL00028 16.07 1.450e-13 548-565
}			BL00028 16.07 2.350e-13 436-453
	1		BL00028 16.07 4.150e-13 492-509
			BL00028 16.07 5.050e-13 296-313
L	l		BL00028 16.07 1.783e-12 520-537

SEQ ID NO:	Accession No.	Description	Results*
	1.10.		BL00028 16.07 3.348e-12 632-649
•		1	BL00028 16.07 5.304e-12 408-425
			BL00028 16.07 5.304e-12 660-677
			BL00028 16.07 4.808e-11 464-481
			BL00028 16.07 7.000e-10 268-285
486	BL00301	GTP-binding elongation	BL00301B 20.09 1.429e-26 128-160
		factors proteins.	BL00301A 12.41 6.400e-15 62-74
487	PD00301	PROTEIN REPEAT MUSCLE CALCIUM-BI.	PD00301B 5.49 7.600e-12 826-837
489	BL00227	Tubulin subunits alpha, beta,	BL00227B 19.29 1.000e-40 52-107
10,		and gamma proteins.	BL00227C 25.48 1.000e-40 113-165
	1		BL00227D 18.46 1.000e-40 222-276
	1		BL00227F 21.16 1.000e-40 382-436
	1		BL00227E 24.15 6.727e-36 326-361
			BL00227A 24.55 2.125e-33 1-35
490	BL00479	Phorbol esters /	BL00479B 12.57 6.625e-09 1271-1287
170.	2200	diacylglycerol binding	
		domain proteins.	
491	BL00479	Phorbol esters /	BL00479B 12.57 6.625e-09 1250-1266
	1 2200	diacylglycerol binding	
		domain proteins.	
492	BL00107	Protein kinases ATP-binding	BL00107A 18.39 5.500e-19 138-169
7/2	5500.07	region proteins.	BL00107B 13.31 1.000e-16 203-219
493	BL50002	Src homology 3 (SH3)	BL50002A 14.19 5.000e-15 392-411
773	DESCUE	domain proteins profile.	BL50002B 15.18 2.500e-09 430-444
494	PR00049	WILM'S TUMOUR	PR00049D 0.00 6.949e-09 87-102
777	1100015	PROTEIN SIGNATURE	
497	BL00914	Syntaxin / epimorphin	BL00914 24.91 6.172e-09 249-299
771	BEOUST	family proteins.	
498	PD00066	PROTEIN ZINC-FINGER	PD00066 13.92 8.200e-16 362-375
770	120000	METAL-BINDI.	PD00066 13.92 4.462e-15 334-347
			PD00066 13.92 8.615e-15 473-486
	1		PD00066 13.92 5.200e-14 306-319
			PD00066 13.92 3.000e-13 390-403
500	PF00780	Domain found in NIK1-like	PF007801 14.69 7.863e-09 293-323
		kinases, mouse citron and	
	1	yeast ROM.	·
501	BL00518	Zinc finger, C3HC4 type	BL00518 12.23 7.333e-09 279-288
		(RING finger), proteins:	
502	DM01418	352 FIBRILLAR	DM01418A 20.83 2.050e-23 1537-1585
302	22.202.120	COLLAGEN CARBOXYL-	DM01418B 22.51 5.895e-21 1632-1674
		TERMINAL.	DM01418C 20.48 8.571e-18 1702-1744
508	BL01052	Calponin family repeat proteins.	BL01052B 15.31 1.000e-09 131-157
512	BL01310	ATPIGI / PLM / MAT8	BL01310 14.74 7.107e-36 27-63
312	BLOISIO	family proteins.	220131011111111111111111111111111111111
515	DM00475	w LOW TRANSPOSASE	DM00475B 12.12 6.019e-09 386-406
212	DW100473	SAPA 12K.	BM001/3B 12:12 0:07,70 07 000 1:0
616	DI 00626	Nt-dnaJ domain proteins.	BL00636A 8.07 5.865e-11 64-81
516	BL00636 PR00625	DNAJ PROTEIN FAMILY	PR00625A 12.84 2.019e-14 76-96
519	FA00023	SIGNATURE	PR00625B 13.48 5.714e-11 106-127
520	DI 00016	Sugar transport proteins.	BL00216B 27.64 6.400e-10 92-142
520	BL00216	Globins profile.	BL01033B 13.81 1.000e-15 38-50
523	BL01033	Src homology 3 (SH3)	BL50002B 15.18 4.750e-12 1075-1089
526·	BL50002	domain proteins profile.	DL30002D 13.10 4./308-12 10/3-1085
531	PR00249	SECRETIN-LIKE GPCR	PR00249G 15.72 8.892e-10 387-409
		SUPERFAMILY	PR00249C 17.08 6.609e-09 223-247
		SIGNATURE	
532	BL00528	Ribosomal protein S4e	BL00528D 27.17 8.012e-09 341-395
		proteins.	

SEQ ID NO:	Accession No.	Description	Results*
534	PR00194	TROPOMYOSIN SIGNATURE	PR00194C 6.38 1.900e-35 109-138 PR00194E 8.74 1.000e-30 195-221 PR00194D 9.57 8.714e-27 139-163 PR00194B 10.24 2.800e-25 84-105 PR00194A 7.86 5.500e-22 48-66
535 .	PR00194	TROPOMYOSIN SIGNATURE	PR00194C 6.38 1.900e-35 109-138 PR00194E 8.74 1.000e-30 195-221 PR00194B 10.24 2.800e-25 84-105 PR00194D 9.57 1.900e-23 139-163 PR00194A 7.86 5.500e-22 48-66
538	PR00019	LEUCINE-RICH REPEAT SIGNATURE	PR00019A 11.19 5.050e-11 110-124
541	BL00540	Ferritin iron-binding regions proteins.	BL00540A 15.06 1.000e-40 32-73 BL00540B 18.82 1.000e-40 123-178 BL00540C 13.00 7.750e-14 188-200
546	PR00153	CYCLOPHILIN PEPTIDYL-PROLYL CIS- TRANS ISOMERASE SIGNATURE	PR00153E 9.10 2.385e-15 121-137
548	BL00115	Eukaryotic RNA polymerase II heptapeptide repeat proteins.	BL00115Z 3.12 8.213e-09 63-112
549	BL01282	BIR repeat proteins.	BL01282B 30.49 2.373e-12 317-356
551	BL00570	Bacterial ring hydroxylating dioxygenases alpha-subunit signa.	BL00570B 19.03 9.357e-09 277-309
553	PD01427	TRANSFERASE METHYLTRANSFERASE BI.	PD01427B 22.45 7.000e-11 127-168
554	PR00048	C2H2-TYPE ZINC FINGER SIGNATURE	PR00048A 10.52 7.632e-11 447-461
555	PD02637	SERUM PARAOXONASE/ARYLES TERASE P.	PD02637A 14.26 1.000e-40 32-87 PD02637G 13.82 1.000e-40 307-355 PD02637D 13.69 6.053e-36 170-218 PD02637B 10.33 8.875e-34 106-141 PD02637E 11.92 8.200e-28 218-249 PD02637C 7.53 3.520e-27 141-170 PD02637F 15.62 9.438e-26 281-307
556	DM00892	3 RETROVIRAL PROTEINASE.	DM00892C 23.55 2.768e-16 474-508
557	BL00039	DEAD-box subfamily ATP-dependent helicases proteins.	BL00039D 21.67 5.179e-36 294-340 BL00039A 18.44 7.955e-29 15-54 BL00039C 15.63 1.300e-16 143-167 BL00039B 19.19 2.465e-12 58-84
558	PR00507	N12 CLASS N6 ADENINE- SPECIFIC DNA METHYLTRANSFERASE SIGNATURE	PR00507B 14.16 8.932e-09 83-98
559	BL00383	Tyrosine specific protein phosphatases proteins.	BL00383E 10.35 8.683e-12 242-253
566	PD00066	PROTEIN ZINC-FINGER METAL-BINDI.	PD00066 13.92 5.500e-13 214-227
572	BL01160	Kinesin light chain repeat proteins.	BL01160B 19.54 4.432e-09 76-130
573	BL00422	Granins proteins.	BL00422C 16.18 4.638e-10 49-77
574	PR00319	BETA G-PROTEIN (TRANSDUCIN) SIGNATURE	PR00319A 15.27 7.911e-10 452-469 PR00319A 15.27 2.180e-09 410-427
577	BL00269	Mammalian defensins proteins.	BL00269C 16.52 6.786e-26 73-102 BL00269A 8.53 2.607e-20 8-28 BL00269B

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SEQ ID NO:	Accession No.	Description	Results*
1.0.	110.		19.17 5.500e-17 35-64
578 [.]	PD02327	GLYCOPROTEIN	PD02327B 19.84 2.241e-11 157-179
	Į.	ANTIGEN PRECURSOR	
		IMMUNOGLO.	
579	BL00596	High potential iron-sulfur	BL00596B 13.07 9.743e-09 273-285
		proteins.	
580	BL00915	Phosphatidylinositol 3- and	BL00915C 22.43 8.147e-32 1015-1054
		4-kinases proteins.	BL00915D 27.02 9.217e-27 1092-1128
			BL00915B 22.78 3.382e-25 918-956
			BL00915A 10.09 5.500e-10 756-768
584	BL00038	Myc-type, 'helix-loop-helix'	BL00038B 16.97 7.488e-09 499-520
	ł	dimerization domain	
		proteins.	
585	BL00795	Involucrin proteins.	BL00795C 17.06 9.200e-09 498-543
586	BL00710	Phosphoglucomutase and	BL00710 12.98 9.100e-17 159-174
	1	phosphomannomutase	1
	-	phosphoserine signa.	DI 00510 10 00 5 714 10 04 40
587	BL00518	Zinc finger, C3HC4 type	BL00518 12.23 5.714e-10 34-43
500	ppocses	(RING finger), proteins. GTP1/OBG GTP-BINDING	PR00326A 8.75 5.979e-14 257-278
588	PR00326	1	FRUUSZOM 8.75 3.9796-14 237-278
		PROTEIN FAMILY SIGNATURE	
501	BL00548	Ribosomal protein S3	BL00548 20.58 7.000e-19 66-96
591	BL00348	proteins.	BL00346 20.36 7.0006-19 00-90
592	BL00478	LIM domain proteins.	BL00478B 14.79 1.250e-12 557-572
392	BL004/8	Livi domani protents.	BL00478B 14.79 1.230c-12 337-372
			BL00478B 14.79 0.000c-12 494-509
594	PR00109	TYROSINE KINASE	PR00109B 12.27 3.681e-13 141-160
J3 4	1 100103	CATALYTIC DOMAIN	11001075 12:27 5:0010-15 141-100
		SIGNATURE	
596	PR00049	WILM'S TUMOUR	PR00049D 0.00 9.063e-12 510-525
3,0	1 100047	PROTEIN SIGNATURE	PR00049D 0.00 8.286e-10 513-528
			PR00049D 0.00 9.000e-10 509-524
			PR00049D 0.00 9.429e-10 511-526
599	BL00232	Cadherins extracellular	BL00232B 32.79 4.750e-40 142-190
		repeat proteins domain	BL00232A 27.72 3.793e-22 48-81
	ļ	proteins.	BL00232B 32.79 1.257e-16 251-299
	l		BL00232C 10.65 5.935e-14 249-267
	İ		BL00232D 16.25 3.368e-13 763-778
			BL00232B 32.79 3.512e-11 366-414
600	DM00215	PROLINE-RICH PROTEIN	DM00215 19.43 9.695e-09 513-546
	nnessa	3.	PEOCEST 10 10 0 100 10 10 100
601	PF00583	Acetyltransferase (GNAT)	PF00583B 10.18 9.100e-10 120-130
(00	PROCES	family.	DD002264 B 75 5 050 11 146 165
602	PR00326	GTP1/OBG GTP-BINDING	PR00326A 8.75 5.950e-11 146-167
		PROTEIN FAMILY	1
604	DI 00210	SIGNATURE	PI 003 10C 17 12 6 0000 10 126 170
604	BL00319	Amyloidogenic glycoprotein extracellular domain	BL00319C 17.12 6.000e-10 136-170
	1	proteins.	
607	BL00239	Receptor tyrosine kinase	BL00239F 28.15 4.717e-25 477-522
007	BLUUZ39	class II proteins.	BL00239F 28.13 4.7176-23 477-322 BL00239E 17.14 5.897e-23 423-473
		ciass ir proteins.	BL00239C 17.14 3.8976-23 423-473 BL00239C 18.75 7.600e-17 372-395
608	PD01066	PROTEIN ZINC FINGER	PD01066 19.43 3.357e-32 10-49
UUO	LDOIDOG	ZINC-FINGER METAL-	1 1001000 17.43 3.33 10-32 10-47
	1	BINDING NU.	
609	PR00449	TRANSFORMING	PR00449A 13.20 4.808e-10 5-27
307	1100449	PROTEIN P21 RAS	PR00449D 10.79 5.636e-09 111-125
		SIGNATURE	1203 1472 10.77 3.0300-07 111-123
610	PF00791	Domain present in ZO-1 and	PF00791C 20.98 2.412e-09 1-40
010	1 44 00 / 71	Politani biocont in 50-1 and	1 1 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2

SEQ ID NO:	Accession No.	Description	Results*
		Unc5-like netrin receptors.	
612	PR00109	TYROSINE KINASE CATALYTIC DOMAIN SIGNATURE	PR00109B 12.27 9.234e-13 487-506
613	BL00678	Trp-Asp (WD) repeat proteins proteins.	BL00678 9.67 1.600e-10 104-115 BL00678 9.67 5.737e-09 62-73 BL00678 9.67 8.105e-09 146-157 BL00678 9.67 8.105e-09 276-287
615	PR00334	HMW KININOGEN SIGNATURE	PR00334B 8.69 5.230e-10 460-484 PR00334B 8.69 1.771e-09 464-488 PR00334B 8.69 2.886e-09 466-490 PR00334B 8.69 8.200e-09 458-482
617	DM00215	PROLINE-RICH PROTEIN 3.	DM00215 19.43 5.881e-09 66-99
618	PF00084	Sushi domain proteins (SCR repeat proteins.	PF00084B 9.45 7.188e-10 539-551 PF00084B 9.45 7.300e-09 600-612
619	PR00169	POTASSIUM CHANNEL SIGNATURE	PR00169A 16.77 4.316e-09 72-92
621	BL00845	CAP-Gly domain proteins.	BL00845 16.43 1.900e-25 321-346 BL00845 16.43 9.325e-22 443-468
622	BL01002	Translationally controlled tumor protein.	BL01002D 18.24 4.706e-26 143-171 BL01002C 21.97 6.143e-26 79-110 BL01002A 13.19 1.360e-24 1-24 BL01002B 7.39 3.118e-14 48-62
624	PR00049	WILM'S TUMOUR PROTEIN SIGNATURE	PR00049D 0.00 8.857e-10 1030-1045
627	PR00011	TYPE III EGF-LIKE SIGNATURE	PR00011A 14.06 4.822e-09 475-494
629	PF00930	Dipeptidyl peptidase IV (DPP IV) N-terminal region.	PF009301 15:96 5.000e-15 656-684 PF00930J 8.78 6.045e-12 708-729
630	PF00930	Dipeptidyl peptidase IV (DPP IV) N-terminal region.	PF009301 15.96 5.000e-15 598-626 PF00930J 8.78 6.045e-12 650-671
631	BL00303	S-100/ICaBP type calcium binding protein.	BL00303B 26.15 1.844e-10 365-402
632	BL00114	Phosphoribosyl pyrophosphate synthetase proteins.	BL00114A 17.22 1.000e-40 54-101 BL00114B 15.90 1.000e-40 107-153 BL00114D 21.45 1.000e-40 208-259 BL00114C 18.22 2.895e-34 167-202 BL00114E 14.48 3.647e-25 293-317
635	BL00870	Chaperonins clpA/B proteins.	BL00870F 8.73 4.833e-36 376-425 BL00870G 8.07 6.553e-27 436-470 BL00870E 17.62 3.333e-16 304-359
639	BL00633	Bromodomain proteins.	BL00633B 13.82 9.775e-13 237-262 BL00633B 13.82 4.750e-11 80-105
641	BL00299	Ubiquitin domain proteins.	BL00299 28.84 7.962e-17 47-99
642	PD02102	SUBUNIT E V-ATPASE VACUOLAR ATP SYNTHASE HYDROL.	PD02102A 16.74 4.176e-10 97-141
643	PD02080	T-CELL GLYCOPROTEIN CD8 CHAIN SURFACE ALPHA PRE.	PD02080D 15.22 6.557e-09 269-306
644	BL01245	RIO1/ZK632.3/MJ0444 family proteins.	BL01245F 18.75 7.805e-14 239-276
646	BL00469	Nucleoside diphosphate kinases proteins.	BL00469 22.22 1.000e-40 41-96
649	PR00217	43 KD POSTSYNAPTIC PROTEIN SIGNATURE	PR00217C 10.91 5.945e-09 91-107
651	PR00326	GTP1/OBG GTP-BINDING PROTEIN FAMILY SIGNATURE	PR00326A 8.75 7.600e-11 629-650

SEQ ID NO:	Accession No.	Description	Results*
652	DM00215	PROLINE-RICH PROTEIN 3.	DM00215 19.43 8.322e-09 227-260
653	PF01298	Transferrin binding protein.	PF01298C 15.13 1.000e-08 413-440
658	PR00443	G-PROTEIN ALPHA SUBUNIT GROUP S SIGNATURE	PR00443A 15.16 9.451e-09 89-105
659	BL00518	Zinc finger, C3HC4 type (RING finger), proteins.	BL00518 12.23 5.714e-10 34-43
663	BL00466	TFIIS zinc ribbon domain proteins.	BL00466 25.88 1.000e-32 294-331
664	PD00567	PROTEIN RNA-BINDING RNA REPEAT HYD.	PD00567B 18.23 3.172e-10 411-425
665	BL00030	Eukaryotic RNA-binding region RNP-1 proteins.	BL00030A 14.39 7.882e-11 10-29
669	PR00124	ATP SYNTHASE C SUBUNIT SIGNATURE	PR00124A 8.81 8.347e-11 117-137
670	PD01234	PROTEIN NUCLEAR BROMODOMAIN TRANS.	PD01234B 15.53 2.500e-10 38-56
671	BL00466	TFIIS zinc ribbon domain proteins.	BL00466 25.88 1.000e-32 219-256
672	BL01282	BIR repeat proteins.	BL01282B 30.49 2.068e-12 298-337
673	BL00455	Putative AMP-binding domain proteins.	BL00455 13.31 4.176e-14 201-217
674	BL01160	Kinesin light chain repeat	BL01160B 19.54 8.703e-10 407-461
		proteins.	BL01160B 19.54 2.373e-09 414-468
675	BL00518	Zinc finger, C3HC4 type (RING finger), proteins.	BL00518 12.23 5.286e-10 326-335
676	BL00518	Zinc finger, C3HC4 type (RING finger), proteins.	BL00518 12.23 5.286e-10 335-344
682	PR00761	BINDIN PRECURSOR SIGNATURE	PR00761E 14.32 4.789e-09 499-518
691	BL00415	Synapsins proteins.	BL00415Q 2.23 2.885e-09 83-119
692	PR00211	GLUTELIN SIGNATURE	PR00211B 0.86 6.167e-09 115-136
694	PR00320	G-PROTEIN BETA WD-40 REPEAT SIGNATURE	PR00320C 13.01 7.300e-09 297-312
696	PD02952	KINASE TRANSFERASE CHOLINE PROTEIN MULTIGENE FAMI.	PD02952C 15.76 5.701e-16 263-293 PD02952B 15.57 7.242e-11 243-257 PD02952A 11.84 9.625e-09 131-159
697	PD00066	PROTEIN ZINC-FINGER METAL-BINDI.	PD00066 13.92 7.231e-15 504-517 PD00066 13.92 5.800e-14 220-233 PD00066 13.92 1.000e-11 248-261 PD00066 13.92 5.696e-11 333-346 PD00066 13.92 2.500e-09 361-374
698 ·	PR00205	CADHERIN SIGNATURE	PR00205B 11.39 6.571e-13 167-185
699	PR00049	WILM'S TUMOUR PROTEIN SIGNATURE	PR00049D 0.00 4.966e-09 50-65 PR00049D 0.00 9.237e-09 74-89
701	PR00988	URIDINE KINASE SIGNATURE	PR00988A 6.39 6.600e-15 98-116 PR00988C 13.64 5.605e-13 175-191 PR00988E 8.27 8.393e-13 245-257 PR00988D 5.95 8.250e-11 231-242 PR00988F 12.23 9.820e-11 267-281 PR00988B 11.60 2.317e-10 128-140
702 .	PR00625	DNAJ PROTEIN FAMILY SIGNATURE	PR00625A 12.84 1.804e-13 22-42 PR00625B 13.48 5.821e-13 53-74
706	PF00023	Ank repeat proteins.	PF00023A 16.03 2.286e-09 209-225
708	BL01212	ATP P2X receptors proteins.	BL01212A 34.89 1.000e-40 43-96
/00	DIMIZIZ	ATT 12A receptors proteins.	BL01212E 24.87 1.000e-40 227-282 BL01212D 11.42 6.700e-25 185-209
			BL01212G 11.86 2.800e-24 310-338 BL01212B 19.25 3.571e-21 129-154

SEQ ID NO:	Accession No.	Description	Results*
10.	1.0.		BL01212C 8.40 1.214c-14 162-173 BL01212F 10.12 4.774e-14 291-302
700	BL00194	Thioredoxin family proteins.	BL00194 12.16 3.455e-17 45-58
709 712	BL00134	Acyltransferases ChoActase	BL00439F 26.22 1.000e-40 418-471
/12	BL00439	/COT/CPT family	BL00439E 19.05 2.440e-24 320-349
	ŀ	proteins.	BL00439B 16.82 1.000e-20 167-189
	1	protests.	BL00439H 18.24 4.600e-20 566-592
		1	BL00439A 9.40 1.237e-15 35-52
		1	BL00439D 13.11 4.545e-15 272-290
			BL00439C 13.53 1.730e-11 248-261
			BL00439G 13.40 9.719e-11 513-524
516	BL00412	Neuromodulin (GAP-43)	BL00412D 16.54 8.990e-09 305-356
716	BL00412	proteins.	
718	BL01271	Sodium:sulfate symporter	BL01271D 25.26 5.979e-32 537-592
		family proteins.	BL01271A 8.06 6.250e-18 131-151
			BL01271C 13.62 7.750e-17 464-486
			BL01271B 12.02 1.563e-16 269-294
719	PF00023	Ank repeat proteins.	PF00023B 14.20 2.500e-10 141-151
	1		PF00023A 16.03 4.000e-10 112-128
721	PF00023	Ank repeat proteins.	PF00023A 16.03 1.750e-10 66-82
	1		PF00023B 14.20 5.500e-09 161-171
			PF00023A 16.03 8.714e-09 363-379
725	PR00019	LEUCINE-RICH REPEAT	PR00019B 11.36 1.500e-11 173-187
123	11.000.5	SIGNATURE	PR00019A 11.19 2.800e-11 314-328
	1	,	PR00019A 11.19 5.050e-11 176-190
			PR00019B 11.36 3.520e-09 311-325
			PR00019B 11.36 4.600e-09 541-555
			PR00019B 11.36 5.320e-09 471-485
			PR00019A 11.19 6.000e-09 544-558
		1	PR00019B 11.36 8.200e-09 242-256
			PR00019B 11.36 9.640e-09 127-141
731	PR00681	RIBOSOMAL PROTEIN SI SIGNATURE	PR00681I 8.81 9.897e-09 600-619
736	PD01066	PROTEIN ZINC FINGER	PD01066 19.43 9.581e-31 8-47
130	1 1001000	ZINC-FINGER METAL-	
	i	BINDING NU.	
739	BL00972	Ubiquitin carboxyl-terminal	BL00972A 11.93 1.587e-13 170-188
139	BEOUT	hydrolases family 2 proteins.	BL00972D 22.55 8.826e-11 590-615
740	BL00972	Ubiquitin carboxyl-terminal	BL00972A 11.93 1.587e-13 170-188
/40	BC00312	hydrolases family 2 proteins.	BL00972D 22.55 8.826e-11 590-615
741	DM01688	2 POLY-IG RECEPTOR.	DM01688G 16.45 6.936e-10 85-117
741	PF00646	F-box domain proteins.	PF00646A 14.37 6.625e-09 50-64
753	BL01168	Ribosomal protein S27e	BL01168 15.74 1.000e-40 20-75
133	PLUITO	proteins.	
756	PD00066	PROTEIN ZINC-FINGER	PD00066 13.92 6.885e-10 127-140
756	PD00000	METAL-BINDI.	
	DD00701	PROTEIN REPEAT	PD00301B 5.49 7.231e-09 1019-1030
757	PD00301	MUSCLE CALCIUM-BI.	15003015 5.47 7.2510 05 1015 100
			BL00712B 12.56 1.000e-40 28-66
761	BL00712	Ribosomal protein S17e	BL00712A 6.23 8.855e-19 2-22
		proteins.	PF00878T 17.51 3.818e-09 799-826
762	PF00878	Cation-independent	11.000/01 17.51 5.0100-05 155-020
1		mannose-6-phosphate	
		receptor repeat proteins.	BL00303A 21.77 9.526e-31 3-40
763	BL00303	S-100/ICaBP type calcium	
		binding protein.	BL00303B 26.15 5.737e-30 50-87
766	BL00018	EF-hand calcium-binding	BL00018 7.41 6.087e-09 237-250
		domain proteins.	
768	BL00221	MIP family proteins.	BL00221D 12.33 6.143e-19 180-195
		•	BL00221C 13.36 1.000e-14 135-152
		•	BL00221E 8.47 3.739e-13 203-214

SEQ ID NO:	Accession No.	Description	Results*
10:	110.		BL00221B 10.22 1.750e-12 63-74 BL00221A 6.39 5.200e-12 16-27
			PF00992A 16.67 8.859e-10 214-249
769	PF00992	Troponin.	BL00720B 16.57 8.297e-15 136-160
770	BL00720	Guanine-nucleotide dissociation stimulators CDC25 family sign.	
77 [PR00883	HIGH MOBILITY GROUP- LIKE NUCLEAR PROTEIN SIGNATURE	PR00883A 6.49 8.920e-09 191-205
772	PD01066	PROTEIN ZINC FINGER ZINC-FINGER METAL- BINDING NU.	PD01066 19.43 6.786e-32 8-47
775	BL00301	GTP-binding elongation factors proteins.	BL00301B 20.09 5.500e-31 90-122 BL00301C 11.73 8.200e-15 423-437 BL00301A 12.41 3.842e-13 9-21
776	PR00453	VON WILLEBRAND FACTOR TYPE A DOMAIN SIGNATURE	PR00453A 12.79 4.892e-12 325-343 PR00453B 14.65 1.614e-10 162-177 PR00453A 12.79 3.152e-10 123-141
779	PR00399	SYNAPTOTAGMIN SIGNATURE	PR00399A 9.52 1.730e-13 145-161 PR00399B 14.27 2.059e-13 160-174 PR00399C 12.82 7.324e-12 216-232 PR00399D 14.48 3.930e-10 236-247 PR00399B 14.27 1.915e-09 291-305
780	BL00115	Eukaryotic RNA polymerase II heptapeptide repeat proteins.	BL00115Z 3.12 8.395e-10 123-172 BL00115Z 3.12 4.375e-09 137-186
783	PD00066	PROTEIN ZINC-FINGER METAL-BINDI.	PD00066 13.92 8.800e-14 165-178 PD00066 13.92 8.800e-14 193-206 PD00066 13.92 5.286e-12 249-262 PD00066 13.92 8.269e-10 221-234
786	PF00975	Thioesterase domain proteins.	PF00975B 10.82 2.688e-12 90-104
788	DM01970	0 kw ZK632.12 YDR313C ENDOSOMAL III.	DM01970B 8.60 9.833e-16 632-645
789	BL00740	MAM domain proteins.	BL00740B 19.76 5.378e-12 174-195 BL00740C 15.93 4.000e-11 684-695
793 ' .	PD00289	PROTEIN SH3 DOMAIN REPEAT PRESYNA.	PD00289 9.97 9.500e-12 102-116
795	BL00038	Myc-type, 'helix-loop-helix' dimerization domain proteins.	BL00038A 13.61 3.400e-09 66-82
800	PD01066	PROTEIN ZINC FINGER ZINC-FINGER METAL- BINDING NU.	PD01066 19.43 5.050e-15 233-272
804	BL00663	Vinculin family talin- binding region proteins.	BL00663G 24.17 1.000e-40 364-414 BL00663K 21.52 9.816e-40 735-790 BL00663I 27.27 4.447e-35 514-568 BL00663I 18.16 3.000e-33 690-727 BL00663L 20.67 9.118e-27 802-838 BL00663F 20.78 2.000e-25 292-333 BL00663F 27.89 1.703e-24 436-489 BL00663C 22.59 2.853e-23 104-159 BL00663B 27.86 4.629e-23 42-96 BL00663D 24.77 3.789e-18 179-226 BL00663A 11.51 2.350e-15 18-39 BL00663E 21.19 9.566e-10 227-265
808	PR00010	TYPE II EGF-LIKE SIGNATURE	PR00010C 11.16 7.545e-10 3968-3979
809	PR00010	TYPE II EGF-LIKE SIGNATURE	PR00010C 11.16 7.545e-10 3882-3893

			
SEQ ID NO:	Accession No.	Description	Results*
810	DM00215	PROLINE-RICH PROTEIN 3.	DM00215 19.43 2.929e-10 163-196
811	PR00205 ·	CADHERIN SIGNATURE	PR00205B 11.39 9.182e-15 243-261 PR00205A 14.73 1.000e-12 168-184 PR00205C 13.65 1.783e-12 503-518 PR00205B 11.39 9.294e-11 463-481
813	PR00456	RIBOSOMAL PROTEIN P2 SIGNATURE	PR00456E 3.06 5.146e-11.313-328 PR00456E 3.06 5.146e-11 314-329 PR00456E 3.06 5.146e-11 315-330 PR00456E 3.06 7.938e-10 312-327 PR00456E 3.06 7.938e-10 316-331
818	BL01071	grpE protein.	BL01071A 24.88 8.277e-21 78-124 BL01071B 18.21 5.286e-15 195-219
826	DM00813	AMINOTRANSFERASES CLASS-V PYRIDOXAL- PHOSPHATE ATTACHMENT SI.	DM00813A 20.30 8.898e-17 231-260
828	BL00415	Synapsins proteins.	BL00415P 2.37 9.814e-09 242-278
830	PF00632	HECT-domain (ubiquitin- transferase).	PF00632C 20.66 5.186e-23 1534-1566 PF00632B 18.45 8.393e-22 1480-1508
831	DM00215	PROLINE-RICH PROTEIN 3.	DM00215 19.43 9.695e-09 117-150
832	PD01066	PROTEIN ZINC FINGER ZINC-FINGER METAL- BINDING NU.	PD01066 19.43 4.231e-33 12-51
834	BL00120	Lipases, serine proteins.	BL00120B 11.37 5.846e-09 1319-1334
836	DM00813	AMINOTRANSFERASES CLASS-V PYRIDOXAL- PHOSPHATE ATTACHMENT SI.	DM00813A 20.30 8.898e-17 38-67
838	PD00289	PROTEIN SH3 DOMAIN REPEAT PRESYNA.	PD00289 9.97 8.000e-12 69-83
840	BL00053	Ribosomal protein S8 proteins.	BL00053B 14.56 1.000e-08 900-918
841	PR00970	ARGININE ADP- RIBOSYLTRANSFERASE SIGNATURE	PR00970D 9.96 3.357e-17 129-146 PR00970A 17.73 8.600e-17 30-52 PR00970E 11.23 6.464e-15 177-193 PR00970B 16.37 2.756e-11 58-77 PR00970C 11.05 9.357e-11 89-104
842	BL00250	TGF-beta family proteins.	BL00250A 21.24 7.120e-25 114-150 BL00250B 27.37 4.774e-18 178-214
846	BL00240	Receptor tyrosine kinase class III proteins.	BL00240B 24.70 7.488e-10 156-180
848	BL01095	Chitinases family 18 proteins.	BL01095B 10.82 5.500e-14 24-36 BL01095C 10.76 7.207e-10 246-258
849	BL00027	'Homeobox' domain proteins.	BL00027 26.43 2.500e-34 300-343
850	PR00318	ALPHA G-PROTEIN (TRANSDUCIN) SIGNATURE	PR00318A 7.84 7.088e-09 188-204
851	PF00969	Class II histocompatibility antigen, beta domain proteins.	PF00969A 22.07 5.846e-29 12-55 PF00969B 9.97 6.211e-25 56-92 PF00969C 27.72 7.324e-16 95-145
852	BL00269	Mammalian defensins proteins.	BL00269B 19.17 6.824e-21 34-63 BL00269A 8.53 6.108e-18 1-21
853	PF00777	Sialyltransferase family.	PF00777B 29.69 8.767e-10 407-450
856	DM00191	w SPAC8A4.04C RESISTANCE SPAC8A4.05C DAUNORUBICIN.	DM00191D 13.94 9.083c-10 100-139

SEQ ID	Accession	Description	Results*
NO:	No.		
857	PR00823	PANCREATIC LIPASE	PR00823A 18.01 3.143e-14 19-37
		SIGNATURE	PR00823C 6.88 6.164e-12 56-69
859	BL00678	Trp-Asp (WD) repeat proteins proteins.	BL00678 9.67 6.684e-09 243-254
860	BL00028	Zinc finger, C2H2 type,	BL00028 16.07 8.650e-13 425-442
		domain proteins.	BL00028 16.07 5.696e-12 508-525
			BL00028 16.07 8.826e-12 564-581
			BL00028 16.07 7.577e-11 201-218
			BL00028 16.07 7.577e-11 536-553
	-		BL00028 16.07 7.923e-11 341-358
			BL00028 16.07 8.615e-11 285-302
		1	BL00028 16.07 1.600e-10 592-609
			BL00028 16.07 2.200e-10 229-246
			BL00028 16.07 3.400e-10 257-274
			BL00028 16.07 6.100e-10 313-330
	Ì		BL00028 16.07 7.000e-10 369-386
			BL00028 16.07 8.200e-10 397-414
			BL00028 16.07 5.114e-09 620-637
864	BL01126	Elongation factor Ts proteins.	BL01126A 18.48 5.011e-10 2637-2680
865	BL00353	HMG1/2 proteins.	BL00353B 11.47 1,330e-13 95-145
		-	BL00353B 11.47 5.692e-11 353-403
866	BL00972	Ubiquitin carboxyl-terminal	BL00972A 11.93 4.600e-18 173-191
	}	hydrolases family 2 proteins.	BL00972D 22.55 1.947e-13 576-601
			BL00972E 20.72 2.038e-11 618-640
867	BL00383	Tyrosine specific protein phosphatases proteins.	BL00383E 10.35 2.756e-12 255-266
872	BL00030	Eukaryotic RNA-binding region RNP-1 proteins.	BL00030B 7.03 5.737e-09 69-79
873	BL00303	S-100/ICaBP type calcium	BL00303B 26.15 4.405e-19 50-87
		binding protein.	BL00303A 21.77 8.765e-15 3-40
874	BL00523	Sulfatases proteins.	BL00523A 13.36 6.500e-17 41-58
		-	BL00523B 8.64 5.909e-15 89-101
			BL00523C 12.64 5.500e-13 140-151
			BL00523D 9.89 9.438e-11 293-305
877	BL00183	Ubiquitin-conjugating enzymes proteins.	BL00183 28.97 1.000e-40 42-90
881	PR00081	GLUCOSE/RIBITOL	PR00081B 10.38 6.727e-11 116-128
		DEHYDROGENASE	PR00081A 10.53 3.106e-10 40-58
		FAMILY SIGNATURE	
882	BL00183	Ubiquitin-conjugating	BL00183 28.97 1.391e-39 50-98
	2233.03	enzymes proteins.	
883	BL00183	Ubiquitin-conjugating	BL00183 28.97 1.391e-39 50-98
	2200703	enzymes proteins.	
888	BL00027	'Homeobox' domain	BL00027 26.43 2.929e-30 232-275
	22002/	proteins.	

Results include Accession number, sub type, eMatrix p-value and the position of the signature.

TABLE 4

TABLE 4				
SEQ ID NO:	Pfam Model	Description	E-value	Pfam Score
445	Rap GAP	Rap/ran-GAP	6.2e-121	415.2
446	zf-C2H2	Zinc finger, C2H2 type	7.4e-65	228.9
452	WD40	WD domain, G-beta repeat	0.00017	28.4
465	WD40	WD domain, G-beta repeat	1.3e-19	78.6
483	COesterase	Carboxylesterases	2.1e-128	440.0
484	hexokinase	Hexokinase	0	2009.4
485	zf-C2H2	Zinc finger, C2H2 type	le-135	464.2
486	GTP EFTU	Elongation factor Tu family	3.2e-125	424.7
487	myosin_head	Myosin head (motor domain)	1.5e-283	955.3
488	Glyco transf 8	Glycosyl transferase family 8	4e-12	53.7
489	tubulin	Tubulin/FtsZ family	3.2e-293	987.5
492	pkinase	Eukaryotic protein kinase domain	7.9e-85	295.2
493	SH3	SH3 domain	1.2e-18	75.4
497	Syntaxin	Syntaxin	0.074	-75.1
498	SCAN	SCAN domain	5.4e-67	236.0
499	F-box	F-box domain	0.0002	28.1
501	FHA	FHA domain	1.7e-13	58.3
502	Collagen	Collagen triple helix repeat (20 copies)	6.5e-197	667.6
507	PH	PH domain	3e-15	59.5
508	СН	Calponin homology (CH) domain	0.0069	16.3
512	ATPIG1_PLM_M AT8	ATPIGI/PLM/MAT8 family	5.7e-31	116.3
516	DnaJ	DnaJ domain	1.4e-24	95.1
519	DnaJ	DnaJ domain	6.8e-26	99.5
522	Glycos_transf_2	Glycosyl transferases	1.2e-13	58.8
523	globin	Globin	4.1e-38	137.3
526	myosin_head	Myosin head (motor domain)	0	1057.8
529	Acetyltransf	Acetyltransferase (GNAT) family	5e-11	50.1
530	MSP_domain	MSP (Major sperm protein) domain	1.7e-16	68.2
531	7tm_2	7 transmembrane receptor (Secretin family)	1.3e-59	211.5
534	Tropomyosin	Tropomyosin	7e-177	553.3
535	Tropomyosin	Tropomyosin	3.1e-173	541.9
538	LRR	Leucine Rich Repeat	2.9e-23	90.7
539	tRNA-synt_1b	tRNA synthetases class I (W and Y)	7.9e-79	275.3
540	PAS	PAS domain	2.8e-05	24.9
541	ferritin	Ferritin	9.9e-116	391.6
546	pro isomerase	Cyclophilin type peptidyl-prolyl cis-tr	3.5e-33	
549	KH-domain	KH domain	0.0004	117.6
551	Glyco_transf 8	Glycosyl transferase family 8	0.0004	27.1
554	zf-C2H2	Zinc finger, C2H2 type	2.6e-22	-47.7
555	Arylesterase	Arylesterase	2.3e-211	87.5
556	G-patch	G-patch domain	2.4e-17	715.6
557	DEAD	DEAD/DEAH box helicase		71.1
558	Methyltransf 4	Putative methyltransferase	8.7e-67	214.2
559	DSPc	Dual specificity phosphatase, catalytic dom	0.0095 4.8e-70	-48.4 246.1
563	IPPT	IPP transferase	670 66	220 /
566	zf-C2H2	Zinc finger, C2H2 type	6.7e-66	232.4
570	RNA_pol_L	RNA polymerases L / 13 to 16 kDa subunit	2.6e-19 0.043	77.6 -12.1
571	Armadillo_seg	Armadillo/beta-catenin-like repeat	96.22	100 /
574	WD40	WD domain, G-beta repeat	8.6e-33	122.4
576	PAP2	PAP2 superfamily	1.1e-65	231.6
377	Defensin_propep	Defensin propeptide	1.2e-19	78.7
578	ig ig	Immunoglobulin domain	3e-25	97.3
580	PI3 PI4 kinase		3.5e-16	57.2
	T T T T VIII T T	Phosphatidylinositol 3- and 4-kinase	6.5e-93	322.1

SEQ ID NO:	Pfam Model	Description	E-value	
585	GBP	Guanylate-binding protein, N-terminal domain	4.3e-165	Sco 548.2
586	PGM_PMM_I	Phosphoglucomutase/phosphomannom utase, alp		4.4
587	zf-C3HC4	Zinc finger, C3HC4 type (RING finger) 1.3e-11	110
588	MMR_HSR1	GTPase of unknown function	5.9e-48	41.9
590	zf-DHHC	DHHC zinc finger domain	1.8e-36	172.7
591	Ribosomal_S3_C	Ribosomal protein S3, C-terminal	1.3e-07	134.6 28.0
592	LIM	LIM domain	4.4e-27	
594	pkinase	Protein kinase domain	3.7e-77	103.4
596	PX	PX domain		269.7
599	Cadherin C term	Cadherin cytoplasmic region	2.2e-17	71.2
600	FHA	FHA domain	3.3e-88	306.5
601	Acetyltransf	Acetyltransferase (GNAT) family	3.4e-20	80.5
604	NAP family	Nucleosome assembly protein (NAP)	3.2e-17	70.6
605	RhoGAP	RhoGAP domain	5.5e-12	46.4
606	Armadillo seg	Armadillo/beta-catenin-like repeat	1e-28	108.9
607	pkinase	Protein kinase domain	0.00022	28.0
608	zf-C2H2	Zinc finger, C2H2 type	5.9e-77	269.1
609	ras	Ras family	5.4e-110	378.8
610	ank	Ank repeat	1.2e-16	52.8
612	pkinase	Protein kinase domain	1.6e-08	41.8
613	WD40	WD domain, G-beta repeat	1.6e-69	244.3
614	UBA	UBA/TS-N domain	4.7e-55	196.3
615	Zip	ZIP Zinc transporter	3.6e-12	53.9
618	sushi	Sushi domain (SCR repeat)	8.1e-59	208.8
619	K_tetra	K+ channel tetramerisation domain	1.3e-58	208.2
621	CAP GLY	CAP-Gly domain	1.3e-19	78.6
622	TCTP	Translationally controlled tumor protein	1.9e-48	174.3
628	UQ con	Ubiquitin-conjugating enzyme	5.2e-109	375.5
529	DPPIV_N_term	Dipeptidyl peptidase IV (DPP IV) N- termi	0.0046 5.1e-07	-43.3 -82.1
630	DPPIV_N_term	Dipeptidyl peptidase IV (DPP IV) N- termi	5.5e-07	-83.2
531	efhand	EF hand	2.3e-14	
532	Pribosyltran	Phosphoribosyl transferase domain	4.3e-37	61.1
35	ank	Ank repeat	1.8e-25	136.7
36	MHCK_EF2_kinas e	MHCK/EF2 kinase domain family	1.2e-12	98.0 5.6
37	DUF221	Domain of unknown function DUF221	1.2e-89	
39	bromodomain	Bromodomain	2.2e-89	311.2
41	ubiquitin	Ubiquitin family	2.2e-29 2.2e-21	106.0
44	RIO1	RIO1/ZK632.3/MJ0444 family	1.1e-07	81.9
46	NDK	Nucleoside diphosphate kinase		-14.9
49	zf-C3HC4	Zinc finger, C3HC4 type (RING finger)	1.1e-52 9.4e-12	188.4
51	ABC tran	ABC transporter	7.9e-84	42.4
54	CUB	CUB domain		291.9
55	MHCK_EF2_kinas e	MHCK/EF2 kinase domain family	3e-30 2.6e-09	113.9 -35.3
59	zf-C3HC4	Zinc finger, C3HC4 type (RING finger)		
61	UvrD-helicase	UvrD/REP helicase	1.3e-11	41.9
63	TFIIS	Transcription factor S-II (TFIIS)	0.078	9.7
64	dsrm		2e-22	87.9
65	птт		4.3e-42	153.3
69	OTU	OTT LIST	0.002	24.8
71	TFIIS	Transposinting Co O Tr Co.	1e-19	78.9
72	zf-C3HC4		2e-22	87.9
73	AMP-binding	Zinc finger, C3HC4 type (RING finger) AMP-binding enzyme	1.5e-05	22.3
<i>'</i> 3 '	AMP-hinding		1.6e-86	

SEQ ID NO:	Pfam Model	Description	E-value	Pfam Score
679	MSP domain	MSP (Major sperm protein) domain	5.4e-18	73.2
680	MSP domain	MSP (Major sperm protein) domain	5.5e-11	49.9
683	RNase PH	3' exoribonuclease family	3e-42	153.8
684	lactamase B	Metallo-beta-lactamase superfamily	0.088	-15.6
686	tRNA anti	OB-fold nucleic acid binding domain	0.031	20.9
690	NHL	NHL repeat	8.2e-18	72.6
691	zf-C3HC4	Zinc finger, C3HC4 type (RING finger)	6.1e-09	33.2
693	WD40	WD domain, G-beta repeat	0.025	21.2
694	WD40	WD domain, G-beta repeat	1.1e-23	92.1
696	Choline kinase	Choline/ethanolamine kinase	1.6e-51	184.6
697	zf-C2H2	Zinc finger, C2H2 type	3.4e-74	259.9
698	cadherin	Cadherin domain	2.2e-05	31.3
701	PRK	Phosphoribulokinase / Uridine kinase family	1.1e-79	278.1
702	DnaJ	DnaJ domain	5e-26	99.9
888	PAX	'Paired box' domain	1.1e-87	304.7

	(AIN, ASE	IDE	CTANT	TEPING	GER			BETA-			UMAN 3AS,			, ZINC	Z.		, ZINC			ZINC Z	_	9
PDB annotation	ASE PDZ DON XIDE SYNTH	INITION PEPT	CYTOKINE, HEMOATTR	JMAJIN GLGF REPE/ EUREXIN,	SE BETA-FIN			REDUCTASE	DIMER		Z DOMAIN, H HPTP1E, PTP-1	DATE OF THE OWNER OWNER O	FINGER/DNA	FINGER/DNA	ADING PROTE	FINGER/DNA	FINGER/DNA	UING PROTE	FINGER/DNA	FINGER/DNA	7.00	INGER/DNA)
PDB	OXIDOREDUCTASE PDZ DOMAIN, NNOS, NITRIC OXIDE SYNTHASE	PEPTIDE RECOGNITION PEPTIDE RECOGNITION, PROTEIN	EOCALIZATION CYTOKINE LCF. CYTOKINE, LYMPHOCYTE CHEMOATTRACTANT	FACTOR, FUZ DOMAIN KINASE HCASK, GLGF REPEAT, DHR; PDZ DOMAIN, NEUREXM, SYNDECAN RECEPTOR CI INTERNA	KINASE OXIDOREDUCTASE BETA-FINGER		MEMBRANE	PROTEIN/OXIDOREDUCTASE BETA-	ringer, hei ekodimer		HYDROLASE PDZ DOMÁIN, HUMÁN PHOSPHATASE, HPTP IB, PTP-BAS, SPECIFICITY 2 OF BINDING	2 1 1 2 2	COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA), ZINC	INGER, DINA-BINDING PROLEIN	COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA), ZINC	ringer, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTFIN		COMPLEX (ZINC FINGER/DNA)
Coumpound	NEURONAL NITRIC OXIDE SYNTHASE, CHAIN: A; HEPTA PEPTIDE: CHAIN: B.	PSD-95; CHAIN: A; CRIPT; CHAIN: B;	INTERLEUKIN 16: CHAIN: NULL:	HCASK/LIN-2 PROTEIN; CHAIN: A, B;	NEURONAL NITRIC OXIDE	SYNTHASE (RESIDUES 1- 130); CHAIN: A;	ALPHA-1 SYNTROPHIN	(RESIDUES 77-171); CHAIN:	OXIDE SYNTHASE	 m	TYROSINE PHOSPHATASE (PTP-BAS, TYPE 1); CHAIN: A;		+	PEPTIDE; CHAIN: A; DUPLEX OLIGONITIE FOTTIDE	z, a,	⊢	PEPTIDE; CHAIN: A; DUPLEX (E, B, C;	┢	_	N: B, C;	
SeqFold score	SYNT	PSD-5	INTERI NULL;	HCAS	NEUR	SYNT 130); (ALPH	(RESI	OX	RESI	(PTP-E		QGSR	OLIGO	BINDI	QGSR	PEPTII	٦		0710	BINDI	QGSR 7
PMF Score	66.0	0.65	0.48	8.0	 - 	, 0.81				į	0.74	L				0.11			80.51			
Verify score	0.39	-0.32	0.58	0.26	0.7	_	0.2			100			0.24		7	-0.03						0.16
PSI- BLAST	1.30E-07	0.00014	1.30E-05	1.10E-11	7.00E-11		5.60E-10			2 805 05	2.00E-03		1.70E-30		7	1.70E-23		00 100	4.20E-29		00 300 7	\neg
End AA	401	401	414	404	410	1	350			414			189		+	748		+	9/7		177	_
Start AA	324	355	329	329	325			_		324			601		137) (CT		101	<u> </u>		10,5	~ ?:
Chain ID	٧	∢		¥	4					V			 ∢					4				
PDB D	1b8q	1be9	1116	1kwa	Iqau		- Agh r			3pdz		+	lal L	_	418	_		lath			lalh	∤
SEQ B B S	445	445		445	445	446				445		╅	6		446	_		446			446	

TABLE 5

PDB annotation	FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGERDNA) COMPLEX (ZINC FINGERDNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CXYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGENDNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
Coumpound	OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE, CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE, CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;
SeqFold score										
PMF		1	-	0.99	<u>-</u>	0.05	0.21	_	0.95	86.0
Verify score		-0.05	-0.09	-0.18	-0.24	0.1	-0.35	0.35	0.38	0.11
PSI- BLAST		6.80E-28	8.50E-28	1.50E-28	1.20E-30	5.10E-20	4.20E-25	1.20E-49	1.70E-38	1.00E-44
End		276	303	331	359	133	189	189	220	276
Start		205	224	252	280	53	81	108	136	201
Chain 10		<	Ą	. ✓	∢	V	4	U	U	ပ
PDB DD		lalh	lalh	lalh	lalh	lalh	lalh	y y	Ime y	1me y
SEQ B B		446	446	446	446	446	446	446	446	446

	PDB	Chain	Start	End	PSI-	Verify	PMF	SeqFold	Coumpound	PDB annotation
- .	Θ.	8	¥₩	¥	BLAST	score	score	score		
- ×	1me y	U	223	303	1.70E-47	0.15	_		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 COYYSTAL STRUCTURE, COMPLEX
- >	1me	U	27	105	5.10E-21	0.08	-0.13	·	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX CINC FINGER/DNA) ZINC COMPLEX CINC FINGER/DNA FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX CZNC FINGER/DNA
	J. J. J. J. J. J. J. J. J. J. J. J. J. J	O.	278	360	1.00E-48			97.55	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
1 .	Jme y	U	279	360	1.00E-48	0.11			DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CEYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
	1me y	υ <u>.</u>		133	5.10E-37	0.19	0.75		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
	1me y	O	08	191	1.00E-48	0.53	-		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
	lme y	ပ	83	190	4.20E-27	-0.13	0.36		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
	ime y	O		105	5.10E-12	0.39	0.19		DNA; CHAIN: A. B. D. E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
	103	Ą	108	192	1.20E-20		-	62.34	TRANSCRIPTION FACTOR	COMPLEX (TRANSCRIPTION

PDB annotation	REGULATION/DNA) TFIIIA, 5S GENE; NMR, TFIIIA, PROTEIN, DNA, TRANSCRIPTION FACTOR, 5S RNA 2 GENE, DNA BINDING PROTEIN, ZINC FINGER, COMPLEX 3 (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATIONDNA) TFILIA; 5S GENE; NMR, TFILIA, PROTEIN, DNA, TRANSCRIPTION FACTOR, 5S RNA 2 GENE, DNA BINDING PROTEIN, ZINC FINGER, COMPLEX 3 (TRANSCRIPTION REGULATIONDNA)	COMPLEX (TRANSCRIPTION REGULATIONDNA) TFIIIA; 5S GENE; NMR, TFIIIA, PROTEIN, DNA, TRANSCRIPTION FACTOR, 5S RNA 2 GENE, DNA BINDING PROTEIN, ZINC FINGER, COMPLEX 3 (TRANSCRIPTION REGULATIONDNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATIONON) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INTIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATIONDNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INTIATION, ZNC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION
Соитроила	IIIA; CHAIN: A; 58 RNA GENE; CHAIN: E, F;	TRANSCRIPTION FACTOR IIIA; CHAIN: A; SS RNA GENE; CHAIN: E, F;	TRANSCRIPTION FACTOR IIIA; CHAIN: A; 58 RNA GENE; CHAIN: E, F;	TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B.C. E. F.
SeqFold score				124.38			
PMF		0.53	0.05		0.94	0.99	0.86
Verify score		0.22	-0.07		0.37	-0.14	0.11
PSI- BLAST		1.20 <u>E-20</u>	1.00E-14	1,70E-36	1,705-36	1.40E-35	3.40E-30
End		189	129	281	257	347	189
Start		109	09	108	109	204	09
Chain		¥	∢	∢	. ∀	∢	₹
PDB		92	9	1476	1466	146	9,14,0
SEQ NO.		446	446	446	446	446	446

					·													·							
PDB annotation	REGULATIONDNA), RNA POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION	REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1;	TRANSCRIPTION INITIALION, INITIATOR ELEMENT, YY1, ZINC 2	FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX TREANSCRIPTION REGIT ATION DNA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION,	INITIATOR ELEMENT, YYI, ZINC2	FINGER PROTEIN, DNA-PROTEIN	RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA) YING-YANG 1;	TRANSCRIPTION INITIATION,	FINGER PROTEIN, DNA-PROTEIN	RECOGNITION, 3 COMPLEX	(TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1:	TRANSCRIPTION INITIATION,	INITIATOR ELEMENT, YY1, ZINC 2	FINGER FROI EIN, DNA-FROI EIN	(TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA) YING-YANG 1;	TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2
Coumpound		TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;		YY I; CHAIN: C; ADENO- ASSOCIATED VIRUS PS	CHAIN: A, B;		YY1; CHAIN: C; ADENO-	ASSOCIATED VIRUS PS INITIATOR FLEMENT DNA:	CHAIN: A, B;			YY1; CHAIN: C; ADENO-	ASSOCIATED VIRUS P5	INITIATOR ELEMENT DNA;	CHAIN: A, B;			ASSOCIATED VIRUS PS	INITIATOR ELEMENT DNA;	CHAIN: A, B;	_		YYI; CHAIN: C; ADENO-	ASSOCIATED VIRUS P5	CHAIN: A, B;
SeqFold score					•						•														
PMF		0.87		0.92			96.0					0.72						-					96.0		
Verify		-0.04		-0.12			0.01					0.01						0.09					-0.01		
PSI- BLAST		8.50E-32		2.80E-32			1.70E-28					1.50E-30						2.80E-35					9.80E-36		
End		229		248	1		220					276						276					331		
Start AA		81		106			116					144						169					198		
Chain ID		∢		O			O					O		_				ပ					O		
80 a		146		lubd			lubd					Inpq						lubd					lubd		
SEQ EQ	2	446		446			446					446						446		_			446		

				· · · · ·		·	
PDB annotation	FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATIONDNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INTIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX	(TRANSCRIPTION RECULATION/DNA) COMPLEX (TRANSCRIPTION RANSCRIPTION INTIATION, INTIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, RECHI ATTONA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX CTRANSCRIPTION REGULATIONDNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INTIATION, INTIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN GECOGNITION, 3 COMPLEX (TRANSCRIPTION PEGILIATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION.
Coumpound		YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YY I; CHAIN: C; ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA;
SeqFold score				105.01		·	
PMF score		0.93			-	0.11	-
Verify score		0.08	-0.08			0.24	0.37
PSI- BLAST		5.10E-32	2.80E-31	9.80E-36	3.40£-34	5.10E-23	5.10E-30
End AA		303	359	360	359	133	161
Start AA		205	221	253	256	27	53
Chain		o	U	O	υ .	U	ပ
PDB TD		Inpd	Iubd	1ubd	Iubd	pqnI	lubd
S e S		446	446	446	446	446	446

SEQ S B S	PDB CD	Chain	Start	End . AA	PSI- BLAST	Verify score	PMF	SeqFold	Coumpound	PDB annotation
									CHAIN: A, B;	INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)
446	1ubd	U	58	681	1.40E-32	0.11	66:0		YYI; CHAIN: C; ADENO ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG I; TRANSCRUPTION INITIATION, INITIATOR ELEMENT, YYI, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)
446	2adr		53	107	1.70E-09	0.04	-0.19		ADRI; CHAIN: NULL;	TRANSCRIPTION REGULATION TRANSCRIPTION REGULATION, ADRI, ZINC FINGER, NMR
446	2gli	¥	109	331	2.80E-42	0.05	0.21		ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
446	2gli	٧	116	275	5.10E-29	0.04	0:11		ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
446	2gli	4	165	359	1.10E-41	0.11	-		ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING . PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
446	2gli	٧	. 195	333	2.80E-42			99.39	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
446	2gli	, 4	207	330	3.40E-30	0.46	1		ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
446	2gli	٧	231	358	3.40E-32	0.27			ZINC FINGER PROTEIN GLI1; CHAIN: A: DNA: CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
446	2gli	¥	32	160	3.40E-27	0.48	0.4		ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA-

							_		_	_		_			_				_										
PDB annotation	BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	CVTORVEI ETONI EI A CTITITA	MEMBRANE SKELETON, SPECTRIN,	CALMODULIN-BINDING, ACTIN-	BINDING, 3 CAPPING PROTEIN,	CALCIUM-BINDING, DUPLICATION, REPEAT. 4 SH3 DOMAIN	STRUCTURAL PROTEIN TWO	REPEATS OF SPECTRIN, ALPHA	HELICAL LINKER REGION, 2.2 TANDEM 3-HELIX COLLED-COLLS	STRUCTURAL PROTEIN	STRUCTURAL PROTEIN TWO	REPEATS OF SPECTRIN, ALPHA	HELICAL LINKER REGION, 22	TANDEM 3-HELIX COILED-COILS,	STRUCTURAL PROTEIN	STRUCTURAL PROTEIN TWO	REPEATS OF SPECTRIN, ALPHA	HELICAL LINKER REGION, 22	TANDEM 3-HELIX COLLED-COLLS,	SIRUCIORAL PROTEIN	STRUCTURAL PROTEIN TWO	HELICAL LINKED BEGION 22	TANDEN STEET IN COURT OF COURT	STRICTURAL PROTEIN	MEMBRANE PROTEIN FOUR HELIX	BUNDLE, ALPHA HELIX	HELIX COLLED COIL, CONTRACTLE
Coumpound		ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	AT DITA CDECOMB DE CITABLE	ALFRA SFECTRIN; CRAIN: NULL;				ALPHA SPECTRIN; CHAIN:	A, B, C;			ALPHA SPECTRIN; CHAIN:	A, B, C;				ALPHA SPECTRIN; CHAIN:	A, B, C;		•		ALPHA SPECTRIN; CHAIN:	A, B, C;			SSO1 PROTEIN; CHAIN: A;	THE LANGUAGE TAREAU TO SELECT THE PERSON OF	AUMAN SKELEI AL MUSCLE ALPHA-ACTININ 2; CHAIN:
SeqFold score																		57.3						•••				37.73	04.40
PMF score		0.93	-	8	60.00				0.23				0.03										0.33				0.29		
Verify score		-0.1	0.31	6	87.0				-0.21				-0.17										-0.0-				-0.32		
PSI- BLAST		2.80E-45	8.50E-31	7. 2007	6.80E-16				1.00E-17				5.60E-05					1.70E-23					1.70E-23				0.00028	1 201	1.705-19
End		278	247	١	139				142				201					762		_			259		_		121	650	707
Start AA		83	88		٠ ر				2				2					37					æ ——	·		•	10	5	77
Chain ED		< .	V						A				A					∀					∢				٧		
PDB CI		2gli	2gli	1	1aj3				lcun				lcun					lcun				1	Ican				1fio		ndor
S B S		446	446	!					447				447					447					447			_	447	447	Ì

PDB annotation	PROTEIN	CONTRACTILE PROTEIN TRIPLE- HELIX COILED COIL, CONTRACTILE PROTEIN	CONTRACTILE PROTEIN TRIPLE. HELIX COILED COIL, CONTRACTILE PROTEIN	CHAPERONE HOP, TPR-DOMAIN, PEPTIDE-COMPLEX, HELICAL REPEAT, HSP90, 2 PROTEIN BINDING	CHAPERONE HOP, TPR-DOMAIN, PEPTIDE-COMPLEX, HELICAL REPEAT, HSP90, 2 PROTEIN BINDING	CHAPERONE HOP, TPR-DOMAIN.	PEPTIDE-COMPLEX, HELICAL	REPEAT, HSC70, 2 HSP70, PROTEIN BINDING	SIGNALING PROTEIN PEROXISMORE	RECEPTOR 1, PTS1-BP, PEROXIN-5,	PTS1 PROTEIN-PEPTIDE COMPLEX,	TETRATRICOPEPTIDE REPEAT, TPR, 2 HELICAL REPEAT	SIGNAT ING PROTEIN PEROXISMORE	RECEPTOR 1, PTS1-BP, PEROXIN-5,	PTSI PROTEIN-PEPTIDE COMPLEX,	TETRATRICOPEPTIDE REPEAT, TPR, 2 HELICAL REPEAT	TRANSCRIPTION INHIBITOR BETA- PROPELLER	METHYLTRANSFERASE	CHEMOTAXIS RECEPTOR	METHYLATION	METHYLTRANSFERASE GNMT, S. ADENOSYL-L-METHIONINE: GI VCINE METHYL TB ANSFERASE	מבו כוועם וווים וווים וויסטים
Coumpound	A;	HUMAN SKELETAL MUSCLE ALPHA-ACTININ 2; CHAIN: A;	HUMAN SKELETAL MUSCLE ALPHA-ACTININ 2; CHAIN: A;	TPR2A-DOMAIN OF HOP; CHAIN: A; HSP90-PEPTIDE MEEVD; CHAIN: B;	TPR2A-DOMAIN OF HOP; CHAIN: A; HSP90-PEPTIDE MEEVD; CHAIN: B;	TPR 1-DOMAIN OF HOP	CHAIN: A, B; HSC70-	PEPTIDE; CHAIN: C, D;	PEROXISOMAL TARGETING	SIGNAL I RECEPTOR;	CHAIN: A, B; PTSI-	CONTAINING PEPTIDE;	DEPOVISONAL TABGETING	SIGNAL 1 RECEPTOR:	CHAIN: A, B; PTS1-	CONTAINING PEPTIDE;	TRANSCRIPTIONAL REPRESSOR TUPI; CHAIN: A, B, C;	CHEMOTAXIS RECEPTOR	CHER: CHAIN: NULL:		GLYCINE N- METHYLTRANSFERASE; CHAIN: A B.	CITATION: A, D,
SeqFold score																						
PMF		0.16	0.11	0.27	0	03	3		0.16				070	Ì			0.19	0.29			0.21	
Verify		-0.31	-0.19	-0.26	-0.49	0.41	;		-0.37				013	71.7			0.01	-0.22			-0.32	
PSI- BLAST		0.0012	1.70E-19	5.10E-05	0.0014	1 20E-08	. 202-702-1		0.0084				4 20E 07	10-707:1			5.10E-38	0.0098			2.80E-06	
End AA		322	265	190	220	186	8		278				480	3			213	146			148	
Start		256	27	110	128	2	.		128				280	· }			4	2			83	
Chain		4	٧	V .	∀	4	¢		4			•		:			4				4	
PDB UD		Iquu	lquu	lelr	lelr	in last	A D		155				- L				lerj	laf7			lxva	
SEQ NO.		447	447	450	450	450	2		450				957	3			452	458			458	

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PDB annotation	LIM DOMAIN CONTAINING PROTEINS LIM DOMAIN CONTAINING PROTEINS, METAL-BINDING PROTEIN, ZINC 2 FINGER	CLATHRIN CLATHRIN, TRISKELION, COATED VESICLES, ENDOCYTOSIS, SELF- 2 ASSEMBLY, ALPHA-ALPHA SUPERHELIX	SIGNALING PROTEIN LIM DOMAIN CONTAINING PROTEINS, METAL- BINDING PROTEIN	TOXIN BINDING PROTEIN TWO DOMAINS: BETA PROPELLER AND ALPHA/BETA FOLD	TOXIN BINDING PROTEIN TWO DOMAINS: BETA PROPELLER AND ALPHA/BETA FOLD	TRANSCRIPTION INHIBITOR BETA- PROPELLER	TRANSCRIPTION INHIBITOR BETA- PROPELLER	TRANSCRIPTION INHIBITOR BETA- PROPELLER	TRANSCRIPTION INHIBITOR BETA- PROPELLER	TRANSCRIPTION INHIBITOR BETA- PROPELLER	COMPLEX (GTP-BINDING/TRANSDUCER) BETA!, TRANSDUCIN BETA SUBUNIT; GAMMAI, TRANSDUCIN GAMMA SUBUNIT; COMPLEX (GTP-BINDING/TRANSDUCER), G PROTEIN, HETEROTRIMER 2 SIGNAL TRANSDUCTION
Coumpound	QCRP2 (LIMI); CHAIN: NULL;	CLATHRIN HEAVY CHAIN; CHAIN: A;	CYSTEINE AND GLYCINE. RICH PROTEIN CRP2; CHAIN: A;	TOLB PROTEIN; CHAIN: A;	TOLB PROTEIN; CHAIN: A;	TRANSCRIPTIONAL REPRESSOR TUP1; CHAIN: A, B. C:	TRANSCRIPTIONAL REPRESSOR TUP1; CHAIN: A, B, C;	TRANSCRIPTIONAL REPRESSOR TUP1; CHAIN: A, B, C;	TRANSCRIPTIONAL REPRESSOR TUP1; CHAIN: A, B, C;	TRANSCRIPTIONAL REPRESSOR TUP1; CHAIN: A, B. C:	GT-ALPHA/GI-ALPHA CHIMERA; CHAIN: A; GT- BETA; CHAIN: B; GT- GAMMA; CHAIN: G;
SeqFold score										•	101.01
PMF	0.09	0.16	0.04	0.77	0.15	0.28	0.76	0.15	0.41	0.01	·
Verify score	-0.46	0.11	-0.39	0.31	0.51	0.31	9.0	0.31	0.36	-0.03	
PSI- BLAST	0.0042	5.10E-33	0.0098	0.0012	0.007	1.70E-48	5.10E-72	6.80E-63	8.50E-64	1.70E-61	1.40E-84
End	879	718	879	481	499	405	200	155	665	639	499
Start AA	851	351	851	225	341	167	181	220	282	330	129
Chain ID		∀	< −	<	<	∢	<	4	¥	¥	æ
PDB CI	la7i	1689	Ica	lerz	lorz	lerj	lerj	lerj	lerj	lerj	lgot
S B S	459	459	459	465	465	465	465	465	465	465	465

			Т		_	_		<u> </u>
PDB annotation	COMPLEX (GTP- BINDING/TRANSDUCER) BETA1, TRANSDUCIN BETA SUBUNIT; GAMMA1, TRANSDUCIN GAMMA SUBUNIT; COMPLEX (GTP- BINDING/TRANSDUCER), G PROTEIN, HETEROTRIMER 2 SIGNAL	COMPLEX (GTP-BINDING/TRANSDUCER) BETA!, TRANSDUCIN BETA SUBUNIT; GAMMA, TRANSDUCIN GAMMA SUBUNIT; COMPLEX (GTP-BINDING/TRANSDUCER), G PROTEIN, HETEROTRIMER 2 SIGNAL TRANSDUCTION	OXIDOREDUCTASE ENZYME, NITRITE REDUCTASE, OXIDOREDUCTASE, DENITRIFICATION, 2 ELECTRON TRANSPORT, PERPLASMIC	TRANSFERASE ESAI HAT, ESAI PROTEIN, ESAIP; HISTONE ACETYLTRANSFERASE, COENZYME A	TRANSFERASE TRANSFERASE (PHOSPHORYL)	TRANSFERASE TRANSFERASE (PHOSPHORYL)		
Соитроина	GT-ALPHA/GI-ALPHA CHIMERA; CHAIN: A; GT- BETA; CHAIN: B; GT- GAMMA; CHAIN: G;	GT-ALPHA/GI-ALPHA CHIMERA; CHAIN: A; GT- BETA; CHAIN: B; GT- GAMMA; CHAIN: G;	CYTOCHROME CDI NITRITE REDUCTASE; CHAIN: A, B;	ESAI HISTONE ACETYLTRANSFERASE; CHAIN: A;	PHOSPHOGLYCERATE MUTASE; CHAIN: A, B;	PHOSPHOGL YCERATE MUTASE; CHAIN: A, B;	TRANSFERASE (PHOSPHORYL) PHOSPHOGLYCERATE MUTASE (E.C.2.7.5.3) DE- PHOSPHO ENZYME 3PGM 4	TRANSFERASE (PHOSPHORYL) PHOSPHOGLYCERATE MUTASE (C.C.2.7.5.3) DE- PHOSPHO ENZYME 3PGM 4
SeqFold score					238.98			226.92
PMF	0.93	0.13	0.37	H .		_	_	
Verify score	0.57	0.28	0.2	0.6		68.0	0.51	
PSI- BLAST	1.40E-84	8.50E-64	4.20E-36	0	6.80E-75	6.80E-75	7.00E-75	7.00E-75
End	499	639	498	290	257	. 652	249	250
Start		326	167	17	16	16	91	16
Chain	æ	m ·	< −	4	V	4		
PDB CD	1got	Igot	1qks	16,7	Iqhf	lqhf	3рg т	3pg m
SEQ NO.	465	465	465	468	470	470	470	470

PDB annotation		TRANSCRIPTION REGULATION PROTO-ONCOGENE, NUCLEAR BODIES (PODS), LEUKEMIA, 2 TRANSCRIPTION REGULATION			LIGASE CBL, UBCH7, ZAP-70, E2, UBIQUITIN, E3, PHOSPHORYLATION, 2 TYROSINE KINASE, UBIQUITINATION, PROTEIN DEGRADATION,	LIGASE CBL, UBCH7, ZAP-70, E2, UBIQUITIN, E3, PHOSPHORYLATION, 2 TYROSINE KINASE, UBIQUITINATION, PROTEIN DEGRADATION,	METAL BINDING PROTEIN RING FINGER PROTEIN MATI; RING FINGER (C3HC4)	METAL BINDING PROTEIN RING FINGER PROTEIN MAT1; RING FINGER (C3HC4)	DNA-BINDING PROTEIN V(D)J RECOMBINATION ACTIVATING PROTEIN I; RAGI, V(D)J RECOMBINATION, ANTIBODY, MAD, RING FINGER, 2 ZINC BINUCLEAR
Coumpound	TRANSFERASE (PHOSPHORYL) PHOSPHOGLYCERATE MUTASE (E.C.2.7.5.3) DE- PHOSPHO ENZYME 3PGM 4	TRANSCRIPTION FACTOR PML; CHAIN: NULL;	VIRUS EQUINE HERPES VIRUS-1 (C3HC4, OR RING DOMAIN) 1CHC 3 (NMR, 1 STRUCTURE) 1CHC 4	VIRUS EQUINE HERPES VIRUS-1 (C3HC4, OR RING DOMAIN) ICHC 3 (NMR, 1 STRUCTURE) ICHC 4	SIGNAL TRANSDUCTION PROTEIN CBL; CHAIN: A; ZAP-70 PEPTIDE; CHAIN: B; UBIQUITIN-CONJUGATING ENZYME E12-18 KDA UBCH7; CHAIN: C;	SIGNAL TRANSDUCTION PROTEIN CBL; CHAIN: A; ZAP-70 PEPTIDE; CHAIN: B; UBIQUITIN-CONJUGATING ENZYME E12-18 KDA UBCH7; CHAIN: C;	CDK-ACTIVATING KINASE ASSEMBLY FACTOR MATI; CHAIN: A:	CDK-ACTIVATING KINASE ASSEMBLY FACTOR MAT1; CHAIN: A:	RAGI; CHAIN: NULL;
SeqFold score									
PMF score		0.04	96.0	99.0	0.51	0.6	0.71	0.43	0.93
Verify score	0.72	-0.13	0.11	-0.45	-0.26	-0.36	-0.15	0.21	-0.05
PSI- BLAST	1.50E-67	3.40E-07	5.60E-12	1.70E-11	8.40E-14	1.70E-11	5.60E-13	3.40E-06	2.80E-25
End	251	70	-73	9/	11	17	73	08	112
Start AA	16	27		29	27	56	28	23	20
Chain ID					. ∢	∢	¥	¥	
PDB ID	3pg m	1bor	Ichc	1chc	1464	1Bv	1825	1825	Irmd
SEQ NO:	470	473	473	473	473	473	473	473	473

PDB annotation	CLUSTER, ZINC FINGER, DNA- BINDING PROTEIN	DNA-BINDING PROTEIN V(D)J RECOMBINATION ACTIVATING PROTEIN I; RAGI, V(D)J RECOMBINATION, ANTIBODY, MAD, RING FINGER, 2 ZINC BINUCLEAR CLUSTER, ZINC FINGER, DNA- BINDING PROTEIN	DNA-BINDING PROTEIN V(D)J RECOMBINATION ACTIVATING PROTEIN 1; RAG1, V(D)J RECOMBINATION, ANTIBODY, MAD, RING FINGER, 2 ZINC BINUCLEAR CLUSTER, ZINC FINGER, DNA- BINDING PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX TRANSCRIPTION REGULATION/DNA)		LIPID TRANSPORT APO A-I; LIPOPROTEIN, LIPID TRANSPORT, CHOLESTEROL METABOLISM, 2 ATHEROSCLEROSIS, HDL, LCAT- ACTIVATION	STRUCTURAL PROTEIN TWO REPEATS OF SPECTRIN, ALPHA HELICAL LINKER REGION, 2.2 TANDEM 3-HELIX COILED-COILS, STRUCTURAL PROTEIN	
Coumpound		RAG1; CHAIN: NULL;	RAGI; CHAIN: NULL;	YYI; CHAIN: C: ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	ZINC FINGER DNA BINDING DOMAIN ZINC-FINGER (ZFY- 6T) (NMR, 13 STRUCTURES) 5ZNF 3	APOLIPOPROTEIN A-1; CHAIN: A, B, C, D;	ALPHA SPECTRIN; CHAIN: A, B, C;	DNA-BINDING HIGH MOBILITY GROUP PROTEIN FRAGMENT-B (HMGB)
SeqFold score			54.37			60.19	51.86	·
PMF		0.86		0.06	0.12			0.01
Verify score		-0.26		-0.51	-0.67			-0.45
PSI- BLAST		5.10E-21	2.80E-25	3,40E-11	0.0042	0.00056	0.00098	0.0014
End		114	114	194	167	206	219	204
Start		23	7	68	140	П	13	164
Chain				ပ		∀	٧	
aga G		1rmd	1rmd	lubd	Sznf	lav1	Icun	1hm e
SEQ SO B		473	473	473	473	475	475	475

PDB annotation		7. 13.	LE CONTRACTILE PROTEIN TRIPLE- HELIX COILED COIL, CONTRACTILE PROTEIN	CHAPERONE HSP40; CHAPERONE, HEAT SHOCK, PROTEIN FOLDING, DNAK	CHAPERONE HSP40; CHAPERONE, HEAT SHOCK, PROTEIN FOLDING, DNAK	CHAPERONE HOP, TPR-DOMAIN, PEPTIDE-COMPLEX, HELICAL REPEAT, HSP90, 2 PROTEIN BINDING	CHAPERONE HÓP, TPR-DOMAIN, PEPTIDE-COMPLEX, HELICAL REPEAT, HSC70, 2 HSP70, PROTEIN BINDING	IG SIGNALING PROTEIN PEROXISMORE RECEPTOR 1, PTS1-BP, PEROXIN-5, PTS1 PROTEIN-PERTIDE COMPLEX, TETRATRICOPEPTIDE REPEAT, TPR, 2 HELICAL REPEAT	MOLECULAR CHAPERONE HDJ-1; MOLECULAR CHAPERONE	MOLECULAR CHAPERONE HDJ-1; MOLECULAR CHAPERONE	MOLECULAR CHAPERONE HDJ-1; MOLECULAR CHAPERONE	CAT OTTO DIAGORA VALORIA
Coumpound	(DNA-BINDING 1HME 3 HMG-BOX DOMAIN B OF RAT HMG1) (NMR, 1 STRUCTURE) 1HME 4	DNA-BINDING HIGH MOBILITY GROUP PROTEIN 1 (FIMG1) BOX 2, COMPLEXED WITH 1HSM 3 MERCAPTOETHANOL (NMR, MINIMIZED AVERAGE STRUCTURE) 1HSM 4	HUMAN SKELETAL MUSCLE ALPHA-ACTININ 2; CHAIN: A;	DNAJ; CHAIN: NULL;	DNAJ; CHAIN: NULL;	TPR2A-DOMAIN OF HOP; CHAIN: A; HSP90-PEPTIDE MEEVD: CHAIN: B:	TPR I-DÓMAIN OF HOP; CHAIN: A, B; HSC70- PEPTIDE; CHAIN: C, D;	PEROXISOMAL TARGETING SIGNAL I RECEPTOR; CHAIN: A, B; PTSI- CONTAINING PEPTIDE; CCHAIN: C, D:	HUMAN HSP40; CHAIN: NULL:	HUMAN HSP40; CHAIN: NULL:	HUMÁN HSP40; CHAIN: NULL;	NET IPOCAT CRI DEL TA.
SeqFold score			55.34	72.19						80.39		
PMF score		0.12				-0.09	0.05	-0.02			_	0.30
Verify score		-0.39				0.38	0.42	ō	1.05		0.98	-0.35
PSI- BLAST		0.00084	4.20E-08	1.20E-19	1.20E-19	1.20E-19	8.50E-24	1.00E-18	1.20E-19	1.40E-30	1.40E-30	0.00013
End		210	222	182	181	129	123	108	180	187	176	76
Start AA		164	-	801	110	9 .	٥	7	107	107	109	43
Chain ID			¥			⋖	∢	∢				4
PDB UD		1hsm	lquu	1bq0	1bq0	lelr	le[w	1fch	1hdj	[þ4]	1hdj	1bif
SEQ NO:		475	475	477	477	477	477	477	477	477	477	479

PDB annotation	BINDING, MYRISTOYLATION, NEURONAL SPECIFIC GUANYLATE 2 CYCLASE ACTIVATOR	SIGNALING PROTEIN CALCIUM BINDING, SIGNALING DOMAIN, NPF BINDING, FW BINDING, 2 EF-HAND, EH DOMAIN, SIGNALING PROTEIN	METAL BINDING PROTEIN CAVP; EF- HAND FAMILY, CALCIUM BINDING PROTEIN, NMR			STRUCTURAL PROTEIN HELIX-TURN-HELIX	METAL TRANSPORT CALMODULIN, HIGH RESOLUTION, DISORDER	METAL BINDING PROTEIN YEAST FREQUENIN EF-HAND, CALCIUM	CALCIUM-BINDING PROTEIN CALCIUM-MYRISTOYL SWITCH, CALCUIM-BINDING PROTEIN	GENE REGULATION POZ DOMAIN;	PROTEIN-PROTEIN INTERACTION	REPRESSOR, ZINC-FINGER PROTEIN,	PROTEIN STRUCTURE,	PROMYELOCYTIC LEUKEMIA, GENE REGULATION	LIGASE CYCLIN A/CDK2- ASSOCIATED PROTEIN P45; CYCLIN A/CDK2-ASSOCIATED PROTEIN P19;
Coumpound	CHAIN: A, B;	EPIDERMAL GROWTH FACTOR RECEPTOR PATHWAY CHAIN: A;	CALCIUM VECTOR PROTEIN; CHAIN: A;	CALCIUM-BINDING PROTEIN CALMODULIN COMPLEXED WITH CALMODULIN-BINDING DOMAIN OF ICDM 3 CALMODULIN-DEPENDENT PROTEIN KINASE II ICDM 4	CALCIUM-BINDING PROTEIN CALMODULIN (VERTEBRATE) ICLL 3	CARDIAC TROPONIN C; CHAIN: A;	CALMODULIN; CHAIN: A;	CALCIUM-BINDING PROTEIN NCS-1; CHAIN: A;	RECOVERIN; CHAIN: NULL;	PROMYELOCYTIC	LEUKEMIA ZINC FINGER				SKP2; CHAIN: A, C, E, G, I, K, M, O; SKP1; CHAIN: B, D, F, H, J, L, N, P;
SeqFold score											•	,			
PMF		0.92	0.42	0.51 0.51	0.13	0.12	0.27	0.21	0.05	0.63					0.69
Verify score		-0.18	-0.3	0.54	0.07	0.28	0.09	0.14	-0.43	0.11					-0.06
PSI- BLAST		0.00014	0.00028	8.40E-05	4.20E-05	0.00017	0.00014	5.60E-05	0.00028	1.70E-24					1.10E-05
End		83	83	107	103	86	103	84	96	143					168
Start AA		41	14	41	14	43	4	43	42	22					52
Chain		4	<	∢		4	4	٧		4					В
PDB ID		1007	1c7w	n n	1cll	146	lexr	1fpw	1 iku	1 buo					1fqv
SEQ NO D		479	479	479	479	479	479	479	479	480					480

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PDB annotation	SKPI, SKP2, F-BOX, LRR, LEUCINE- RICH REPEAT, SCF, UBIQUITIN, 2 E3, UBIQUITIN PROTEIN LIGASE	LIGASE SKP2 F-BOX; SKP1; SKP1, SKP2, F-BOX, LRR, LEUCINE-RICH REPEAT, SCF, UBIQUITIN, 2 E3, UBIQUITIN PROTEIN LIGASE	LIGASE CYCLIN A/CDK2- ASSOCIATED P45; CYCLIN A/CDK2- ASSOCIATED P19; SKP1, SKP2, F-BOX, LRK3, LEUCINE-RICH REPEATS, SCF, 2 UBIQUITIN, E3, UBIQUITIN PROTEIN LIGASE	·				INSECT IMMUNITY INSECT IMMUNITY, LPS-BINDING, HOMOPHILIC ADHESION	GROWTH FACTOR/GROWTH FACTOR RECEPTOR FGF, FGFR, IMMUNOGLOBULN-LIKE, SIGNAL TRANSDUCTION, 2 DIMERIZATION, GROWTH FACTOR/GROWTH FACTOR	VIRUSVIRAL PROTEIN, RECEPTOR CD155, PVR, HUMAN POLIOVIRUS, ELECTRON MICROSCOPY, 2 POLIOVIRUS-RECEPTOR COMPLEX, VIRUSVIRAL PROTEIN PROFEDIOR	GROWTH FACTOR/GROWTH FACTOR RECEPTOR FGF2; FGFR2; IMMUNOGLOBULIN (IG)LIKE
Coumpound		CYCLIN A/CDK2- ASSOCIATED P19; CHAIN: A, C; CYCLIN A/CDK2- ASSOCIATED P45; CHAIN: B, D;	SKP2; CHAIN: A, C; SKP1; CHAIN: B, D;	OXIDOREDUCTASE(OXYGE N(A)) GALACTOSE OXIDASE (E.C.1.1.3.9) (PH 4.5) 1GOF 3	OXIDOREDUCTASE(OXYGE N(A)) GALACTOSE OXIDASE (E.C.1.1.3.9) (PH 4.5) 1GOF 3	OXIDOREDUCTASE(OXYGE N(A)) GALACTOSE OXIDASE (E.C.1.1.3.9) (PH 4.5) 1GOF 3		HEMOLIN; CHAIN: A, B;	FIBROBLAST GROWTH FACTOR 2, CHAIN: A, B; FIBROBLAST GROWTH FACTOR RECEPTOR 1; CHAIN: C, D;	POLIOVIRUS RECEPTOR; CHAIN: R; VP1; CHAIN: 1; VP2; CHAIN: 2; VP3; CHAIN: 3; VP4; CHAIN: 4;	FIBROBLAST GROWTH FACTOR 2; CHAIN: A, B, C, D; FIBROBLAST GROWTH
SeqFold score		·									
PMF		0.96	0.86	-0.08	0.22	-0.18		-0.17	0.21	-0.14	0.17
Verify score		0.48	0.26	0.22	0.31	0.07		0	-0.04	0.03	0.05
PSI- BLAST		4.20E-05	0.00042	1.40E-31	8.50E-13	2.80E-14		1.70E-29	1.40E-12	1.70E-14	5.10E-13
End		141	164	577	588	592		396	396	381	392
Start		52	22	318	346	362		66	303	96	303
Chain D		æ	æ					Y	Ω	ಜ	म
PDB CI		វិទិវ	1fs2	lgof	1gof	1gof		15ih	lcvs	Idgi	lev2
SEQ EQ		480	480	480	480	480		481	481	481	481

PDB annotation	DOMAINS BELONGING TO THE I-SET 2 SUBGROUP WITHIN IG-LIKE DOMAINS, B-TREFOIL FOLD	GROWTH FACTOR/GROWTH FACTOR	MACEFIOR FORZ; FORZ;	DOMAINS BELONGING TO THE I-SET	2 SUBGROUP WITHIN IG-LIKE	DOMAINS, B-TREFOIL FOLD	GROWTH FACTOR/GROWTH FACTOR	IMMUNOGLOBULIN (IG) LIKE	DOMAINS BELONGING TO THE I-SET	2 SUBGROUP WITHIN IG-LIKE DOMAINS, B-TREFOIL FOLD	IMMUNE SYSTEM FC-EPSILON RI-	ALPHA; IMMUNOGLOBULIN FOLD,	GLYCOPKOIEIN, KECEPIOK, 1GE-	CONTRACTILE PROTEIN	IMMUNOGLOBULIN FOLD, BETA	KINASE KINASE, TWITCHIN,	IMMINE SVSTEM CD32: BECEBTOR	IMMUNE SYSTEM CD32, NECETION, FC, CD32, IMMUNE SYSTEM	CHAPERONE HOP, TPR-DOMAIN, PEPTIDE-COMPLEX, HELICAL REPEAT, HSP90, 2 PROTEIN BINDING	HALOPEROXIDASE	HALOPEROXIDASE F;	OXIDOREDUCTASE, PROPIONATE	COMPLEX	HYDROLASE BPHD; HYDROLASE, PCB DEGRADATION		HYDROLASE PNB ESTERASE; ALPHA- BETA HYDROLASE, DIRECTED
Coumpound	FACTOR RECEPTOR 2; CHAIN: E, F, G, H;	FIBROBLAST GROWTH	FACTOR 2; CHAIN: A, B, C, D;	FACTOR RECEPTOR 2;	CHAIN: E, F, G, H;		FIBROBLAST GROWTH	FIBROBLAST GROWTH	FACTOR RECEPTOR 1;	CHAIN: C, D;	HIGH AFFINITY	IMMUNOGLOBULIN	EFSILON RECEPTOR CHAIN: A:	TELOKIN; CHAIN: A		TWITCHIN; CHAIN: NULL;	EC GAMMA BIIB: CHAIN: A:	ו כ טעאנואם הנום, כוזהנוא. ה,	 TPR2A-DOMAIN OF HOP; CHAIN: A; HSP90-PEPTIDE MEEVD; CHAIN: B;	CHLOROPEROXIDASE F;	CHAIN: NOLL;			2-HYDROXY-6-OXO-6- PHENYLHEXA-2.4-	DIENOATE CHAIN: A;	PARA-NITROBENZYL ESTERASE; CHAIN: A;
SeqFold score																			,							
PMF score		0.15					0.1				0.04			-0.07		0.03	100	100	60.0	0.04				0.47		1
Verify score		-0.2					0.02				-0.2			0.13		90.0	70.05	3	-0.06	0.45				0.29		0.39
PSI- BLAST		5.10E-13					1.40E-12				1.70E-16			3.40E-17		8.50E-16	1 70F-14	11-20/	0.00014	0.0017				0.00012		3.40E-93
End		392					396				386			381		382	384	3	363	296				331		606
Start AA		303					303				192			289		292	5	:	281	152				178		43
Chain		O					ပ				 			4			A		₹					∢		∢ .
PDB ID		lev2					levt				1229			1fhg		lkoa	2fch		lefr	1a8s				1c4x		1c7j
S B SE		481					481				481			481		481	481	<u>:</u>	482	483				483		483

PDB annotation	EVOLUTION, ORGANIC ACTIVITY, 2 PNB ESTERASE	LIPASE ESTERASE, SUBSTRATEPRODUCT-BOUND ICLE 9	LIPASE ESTERASE, SUBSTRATEPRODUCT-BOUND ICLE 9	HYDROLASE LINB, 1,3,4,6- TETRACHLORO-1,4- CYCLOHEXADIENE DEHALOGENASE, LINDANE, BIODEGRADATION, ALPHABETA-HYDROLASE	HYDROLASE (SERINE ESTERASE) HYDROLASE (SERINE ESTERASE), HYDROLASE, SERINE ESTERASE, 2 SYNAPSE, MEMBRANE, NERVE, MUSCLE, SIGNAL, NEUROTRANSMITTER 3 DEGRADATION, GLYCOPROTEIN, GPI-ANCHOR, ALTERNATIVE SPLICING	CHOLINESTERASE SERINE HYDROLASE, NEUROTRANSMITTER CLEAVAGE, CATALYTIC 2 TRIAD, ALPHA/BETA HYDROLASE	HYDROLASE ALPHA/BETA HYDROLASE FOLD	HYDROLASE ALPHA/BETA HYDROLASE FOLD	HYDROLASE BILE SALT ACTIVATED LIPASE, ESTERASE, CATALYTIC DOMAIN	SERINE HYDROLASE SERINE HYDROLASE, DEGRADATION OF BREFELDIN A, ALPHA/BETA 2 HYDROLASE FAMILY	
Соимроипа		CHOLESTEROL ESTERASE; ICLE 4 CHAIN: A, B; ICLE 5	CHOLESTEROL ESTERASE; 1CLE 4 CHAIN: A, B; 1CLE 5	HALOALKANE DEHALOGENASE; CHAIN: A;	ACETYLCHOLINESTERASE; CHAIN: A;	ACETYLCHOLINESTERASE; CHAIN: A;	SERINE HYDROLASE; CHAIN: A;	SERINE HYDROLASE; CHAIN: A;	BILE SALT ACTIVATED LIPASE; CHAIN: A;	BREFELDIN A ESTERASE; CHAIN: A, B;	HYDROLASE LPASE (E.C.3.1.1.3) (TRIACYLGLYCEROL
SeqFold score		181.19									179.92
PMF			_	0.57	M.	-	0.58	0.42	-	0.1	
Verify score			0.21	9.6	0.61	0.59	0.17	-0.05	0.5	-0.05	·
PSI- BLAST		1.20E-73	1.20E-73	0.0084	0	0	3.40E-28	5.60E-39	0	5.10E-20	1.20E-71
End		581	593	379	611	612	346	283	612	334	1881
Start		42	89	163		40	142	73 ₁	44	83	42
Chain TO		ď	V .	∀	₽.	٧	٧	A	∀	¥	
PDB CD		Icle	lcle	16v2	1dx4	lea5	levq	levq	1f6w	1.jkm	Прр
SEQ NO D		483	483	483	483	483	483	483	483	483	483

· 🛏 🗆	PDB CI	Chain	Start AA	End	PSI- BLAST	Verify score	PMF score	SeqFold	Coumpound	PDB annotation
1									ILPP 3 HEXADECANESULFONATE ILPP 4 ILPP 71	
1	1lpp		89	593	1.20E-71	0.22	-		HYDROLÁSE LIPASE (E.C.3.1.1.3) (TRIÁCYLGLÝCEROL I PASE) COMPI.EXED WITH	
									ILPP 3 HEXADECANESULFONATE ILPP 4 ILPP 71	
=	Imaa A		38	612	0			368.25	ACETYLCHOLINESTERASE; CHAIN: A, B, C, D;	HYDROLASE MACHE; HYDROLASE, SERINE ESTERASE, ACETYLCHOLINESTERASE, TETRAMER, 2 HYDROLASE FOLD,
15	Imaa A		38	612	0	0.72			ACETYLCHOLNESTERASE; CHAIN: A, B, C, D;	HYDROLASE MACHE; HYDROLASE, SERINE ESTERASE, ACETYLCHOLINESTERASE, TETRAMER, 2 HYDROLASE FOLD, GLYCOSYLATED PROTEIN
15.	1qe3 A		40	602	1.70E-89			242.59	PARA-NITROBENZYL ESTERASE; CHAIN: A;	HYDROLASE PNB ESTERASE; ALPHA- BETA HYDROLASE DIRECTED EVOLUTION
13	Iqe3 A		£4	599	1.70E-89	0.33	-		PARA-NITROBENZYL ESTERASE; CHAIN: A;	HYDROLASE PNB ESTERASE, ALPHABETA HYDROLASE DIRECTED EVOLUTION
I≍	1qfm A		26	397	5.60E-57	0.15	0.11		PROLYL OLIGOPEPTIDASE; CHAIN: A;	HYDROLASE PROLYT. ENDOPEPTIDASE, POST-PROLINE CLEAVING PROLYL OLIGOPEPTIDASE, AMNESIA, ALPHABETA-HYDROLASE, BETA- 2 PROPELLER
Ι Ξ	A A P		87	350	1.20E-35	0	0.03		PROLYL OLIGOPEPTIDASE; CHAIN: A:	HYDROLASE PROLYL ENDOPEPTIDASE, POST-PROLINE CLEAVING PROLYL OLIGOPEPTIDASE, AMNESIA, ALPHA/BETA-HYDROLASE, BETA- 2 PROPELLER
\ =	1thg		46	280	5.10E-80			210.1	HYDROLASE(CARBOXYLIC ESTERASE) LIPASE (E.C.3.1.1.3)	

										·	
PDB annotation			HYDROLASE BILE SALT ACTIVATED LIPASE, BILE SALT STIMULATED HYDROLASE, SERINE ESTERASE, LIPASE	HYDROLASE BILE SALT ACTIVATED LIPASE, BILE SALT STIMULATED HYDROLASE, SERINE ESTERASE, LIPASE	HEXOKINASE ATP/.D-HEXOSE-6- PHOSPHOTRANSFERASE; HEXOKINASE, PHOSPHOTRANSFERASE	HEXOKINASE ATP/.D-HEXOSE-6- PHOSPHOTRANSFERASE; HEXOKINASE, PHOSPHOTRANSFERASE	TRANSFERASE STRUCTURALLY HOMOLOGOUS DOMAINS	COMPLEX (ZINC FINGENDNA) COMPLEX (ZINC FINGENDNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGERDNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA
Coumpound	TRIACYLGLYCEROL HYDROLASE 1THG 3	HYDROLASE(CARBOXYLIC ESTERASE) LIPASE EC.3.1.1.3) TRIACYLGI.YCEROL HYDROLASE ITHG 3	CHOLESTEROL ESTERASE; CHAIN: NULL;	CHOLESTEROL ESTERASE; CHAIN: NULL;	HEXOKINASE; CHAIN: A, B;	HEXOKINASE; CHAIN: A, B;	HEXOKINASE TYPE I; CHAIN: N;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE: CHAIN: B. C:	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER
SeqFold score			318.91			179.92					
PMF		-		1	-			0.11	0.3	0.94	0.11
Verify score	•	0.49	·	0.54	1.19		1.21	-0.49	-0.36	0.16	-0.34
PSI- BLAST		5.10E-80	0	0	0	0	0	1.70E-18	1.70E-23	6.80E-24	1.70E-30
End		583	618	612	910	116	913	204	232	260	204
Start AA		47	39	44	-	-	91	127	152	081	126
Chain					¥	4	z	∢	⋖	∢	O
PDB UI		lthg	2bce	2bce	1bg3	1bg3	Icza	lalh	laih	lalh	Jme y
SEQ B B		483	483	483	484	484	484		485	485	485

														_						_	_		_		.,				_			_
PDB annotation	INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA	INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX	COMPLEY (21NC FINGER/DNA) ZINC	FINGER, PROTEIN-DNA	INTERACTION, PROTEIN DESIGN, 2	CRYSTAL STRUCTORE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC	FINGER, PROJEIN-DNA	COVETAL STRUIGHTER COMPLEX	(ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC	FINGER, PROTEIN-DNA	INTERACTION, PROTEIN DESIGN, 2	CRYSTAL STRUCTURE, COMPLEX	(ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC	FINGER, PROTEIN-UNA	NTERACTION, PROTEIN DESIGN, 2	(ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC	FINGER, PROTEIN-DNA	INTERACTION, PROTEIN DESIGN, 2	CRYSTAL STRUCTURE, COMPLEX	COMPLEX (21NC FINGER/DNA) 21NC	FINGER PROTEIN-DNA	INTERACTION, PROTEIN DESIGN, 2	CRYSTAL STRUCTURE, COMPLEX	(ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC	FINGER, PROTEIN-DNA	INTERACTION, PROTEIN DESIGNAL
Conmpound	PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;	Date: Citable & D. D.	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;		DNA; CHAIN: A, B, D, E;	CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;		DNA; CHAIN: A, B, D, E;	CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;			DNA; CHAIN: A, B, D, E,	CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;	-	DNA; CHAIN: A, B, D, E;	CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;		DNA: CHAIN: A B D E.	CONSENSITS ZINC FINGER	PROTEIN; CHAIN: C, F, G;			DNA; CHAIN: A, B, D, E;	CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, C;
SeqFold score																											•					
PMF score		69.0		8	66:0			_				L					-				_				_	•				_		
Verify score		-0.18		3	-0.0 -0.0		_	0.05				0.27					0.25		_		0.25				130) 				29.0		
PSI- BLAST		5.10E-38		.,	8.50E-41			1.20E-42			-	1.40E-43		_			2.80E-47				5.10E-47				1 SAE 48					2.80E-51		
End	i	232			260			288				316		_			372				372		_		907	}		_		400		
Start		151		-	179			207				235					291				291				310	<u>}</u>				319	_	
Chain		o.			ပ			ပ				ပ					၁				ပ				c	,				၁		
PDB		1me	`		lme ;	`		Ime	<u>~</u>			Ime	×.				Ime	>			1me	^			120	} } }	<u> </u>			lmc	۸	
SEQ 5.0	Ž	485			485			485				485			·		485				485				707	}				485		

PDB annotation	CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA	INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (77NC FINGER MAX)	COMPLEX (ZINC FINGER/DNA) ZINC	FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2	CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC	INTERACTION, PROTEIN DESIGN, 2	CRYSTAL STRUCTURE, COMPLEX	COMPLEX (ZINC FINGER/DNA) ZINC	FINGER, PROTEIN-DNA	INTERACTION, PROTEIN DESIGN, 2	CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC	FINGER, PROTEIN-DNA	INTERACTION, PROTEIN DESIGN, 2	CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC	FINGER, PROTEIN-DNA	INTERACTION, PROTEIN DESIGN, 2	CRYSTAL STRUCTURE, COMPLEX	COMPLEX (ZINC FINGER (DNA) ZINC	FINGER PROTEIN-DNA	INTERACTION, PROTEIN DESIGN, 2	CRYSTAL STRUCTURE, COMPLEX	(ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC	FINGER, PROTEIN-DNA	INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX
Coumpound		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E;	CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;		DNA; CHAIN: A, B, D, E;	PROTEIN; CHAIN: C, F, G;		DNA; CHAIN: A, B, D, E;	CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;		DNA; CHAIN: A, B, D, E;	CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;		DNA; CHAIN: A. B, D, E:	CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;		DNA: CHAIN: A B D E.	CONSENSITION FINGER	PROTEIN; CHAIN: C. F. G.		!	DNA; CHAIN; A, B, D, E;	CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;
SeqFold score										113.26																		•	
PMF score		-	•	_			1	•						_				_				-					1		
Verify score		0.48		69.0			69.0							0.41				0.05				0.02	!				0.1		
PSI- BLAST		5.10E-49		3.40E-49			4.20E-50			4.20E-50				1.00E-49				1.70E-50				1.50E-50					8.50E-51		
End		428		456			456			457				484				212				540					268		
Start		347		375			375			375				403				431				459					487		
Chain TD		. o		C			၁			၁				O				ပ				U					ပ		
PDB ID		lme y		Ime	<u>></u>		lme	<u> </u>		Ime	>			Ime	>			Ime	>			1	^			┪	j E	>	
SEQ B B S		485		485			485			485				485				485				485					485		

													_		_				_		_			_		_				_	_			
PDB annotation	(ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA	INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRITCTURE, COMPLEX	(ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC	FINGER, PROTEIN-DNA	INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRICTLIRE, COMPLEX	(ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC	FINGER, PROTEIN-DNA	INTERACTION, PROTEIN DESIGN, 2	CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC	FINGER, PROTEIN-DNA	INTERACTION, PROTEIN DESIGN, 2	CRYSTAL STRUCTURE, COMPLEX	(ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC	FINGER, PROTEIN-UNA	INTERACTION, PROTEIN DESIGN, 2	CRYSTAL STRUCTORE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC	FINGER, PROTEIN-DNA	INTERACTION, PROTEIN DESIGN, 2	CRYSTAL STRUCTURE, COMPLEX	CLINC FINGENDINA)	COMPLEX (IRANSCRIPTION PEGIT ATTOMONA) COMPLEX	TRANSCRIPTION	REGULATION/DNA), RNA	POLYMERASE III, 2 TRANSCRIPTION	INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION	KEGULA HON/DNA) COMPLEX	REGULATION/DNA). RNA
Coumpound		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;		DNA; CHAIN: A, B, D, E;	CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;		DNA; CHAIN: A, B, D, E;	CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;		DNA: CHAIN: A. B. D. E.	CONSENSUS ZINC FINGER	PROTEIN, CHAIN: C, F, G,		!	DNA; CHAIN: A. B, D, E;	CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G:		DNA; CHAIN: A, B, D, E;	CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;		Co C . ACT AND T AND COM	TFIIIA; CHAIN: A, D; 5S	CHAIN B C E F				TFIIIA; CHAIN: A, D; 5S	KIBOSOMAL KNA GENE;	CITATION 15, C, L, T,
SeqFold score										•																								
PMF		_			1								-					1				0.48					0.76					0.81		
Verify score		0.02			0.03				0.28				0.15	}				0.36				-0.38				3	-0.28					-0.2		
PSI- BLAST		1.50E-50			3.40E-50				1.70E-50				3 40E-50					1.00E-32				5.10E-07				90 207 0	3.40E-27					3.40E-31		
End AA		965			624				652				680	3		_		683				176				1	274					297		
Start		\$15			543	!			571				\$00					627				149					130	`				152		
Chain ID		υ			ပ				O				ļ	,				O				0		_			⋖					V		
PDB		Ime v	,		Ime	٨			Ime	^			1	}	`			Ime	>			Ime	>				1466					146		
SEQ B	į	485			485	!			485				185	}				485				485					485					485		

PDB annotation	POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX	(TRANSCRIPTION REGULATION/DNA), RNA	POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION	REGULATION/DNA) COMPLEX	REGULATION/DNA), RNA	POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION	REGULATION/DNA) COMPLEX	(IRANSCRIPTION RNA REGILI ATION/DNA) RNA	POLYMERASE III, 2 TRANSCRIPTION	INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION	KEGULATION/DNA) COMPLEX	BEGIN ATION/DNA) RNA	POLYMERASE III, 2 TRANSCRIPTION	INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION	REGULATION/DNA) COMPLEX	(IKANSCKIPTION	POLYMERASE III. 2 TRANSCRIPTION	INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION	REGULATION/DNA) YING-YANG 1;	INITIATOR FLEMENT YYL ZINC2	FINGER PROTEIN, DNA-PROTEIN	RECOGNITION, 3 COMPLEX	COMPLEX (TRANSCRIPTION REGULATIONDNA) YING-YANG 1;
Coumpound		TFIIIA; CHAIN: A. D. SS RIBOSOMAL RNA GENE;	CHAIN: B, C, E, F;		TFIIIA; CHAIN: A, D; 5S	RIBOSOMAL RNA GENE;	Chair. B, C, E, I,		TFIIIA; CHAIN: A, D; 5S	RIBOSOMAL RNA GENE;	CHAIN: B, C, E, F;			TFIIIA; CHAIN: A, D; 5S	RIBOSOMAL KNA GENE;	CHAIN: B, C, E, F;			TFIIIA; CHAIN: A, D; 5S	RIBOSOMAL RNA GENE;	CHAIN: B, C, E, F;		,	YY1; CHAIN: C; ADENO-	ASSOCIATED VIRUS PS	CHAIN: A B.			YY1; CHAIN: C; ADENO-ASSOCIATED VIRUS P5
SeqFold score					120.46			,							•														
PMF		-				-			_					0.99					√ 66.0					0.12					0.72
Verify score		0.07							0.11					0.01					0.1	_				-0.22					-0.27
PSI- BLAST		6.80E-34			5.10E-37				5.10E-37					3.40E-37					1.40E-36					5.10E-25					8.50E-27
End		353			484				521					633					682					232					260
Start		208			319				376			_		488					544					127					159
Chain		V			A	:			A		_			Ą					A					ပ					O
PDB TD		1116			¥.]			1466					1466					146					Iubd					lubd
SEQ	Ž	485			485	}			485					485					485					485					485

PDB annotation	TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATIONIDNA)	COMPLEX (TRANSCRIPTION REGULATIONDNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRPTION REGULATIONDNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN TRECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX CTRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION
Coumpound	INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YY1; CHAIN: C; ADENO-
SeqFold							
PMF		0.86	0.93		0.96	-	
Verify		-0.04	0.32	0.18	0.02	0.14	0.63
PSI- BLAST		3.40E-29	1.20E-29	8.50E-32	4.20E-46	7.00E-52	9.80E-59
End AA		288	316	344	345	372	400
Start		182	215	238	240	268	296
Chain 10		U	U	SO.	U	o .	ပ
PDB		1 ubd	Iubd	1ubd	lubd	1ubd	Jubd
SEQ NO NO		485	485	485	485	485	485

PDB annotation	REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATIONDNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATIONDIA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION, REGULATIONDIA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)
Coumpound	ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B:
SeqFold score			100.22		,	
PMF score		-		-		1
Verify		. 0.04		0.26	0.1	0.09
PSI- BLAST		2.80E-59	2.80E-59	2.80E-55	8.40E-56	4.20E-59
End		456	457	512	296	652
Start		346	349	401	485	541
Chain ID		U	U	ပ	U	U
EDB CD		pqn1	1ubd	Iubd	Inpq	lubd
SEQ Se Se		485	485	485	485	485

PDB annotation	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INTIATION, INTIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATIONDAA)	: <u>;</u>	II; COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	JI; COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	LII; COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	ë.	II. COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLJ; GLJ, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	-	Ë	-
Coumpound	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A: DNA: CHAIN: C.
SeqFold						110.36				
PMF score	0.98	0.09	0.47	-	-		-i	0.88		86.0
Verify score	0.06	-0.05	0.01	9.4	0.53		0.54	0.01	-0.12	0.09
PSI- BLAST	3.40E-34	1.00E-26	5.60E-44	3.40E-31	1.40E-68	2.80E-77	2.80E-77	1.40E-73	1.10E-73	1.40E-72
End	089	262	346	343	402	458	458	542	654	681
Start AA	579	131		207	263	349	319	375	487	516
Chain TD	ပ	¥	¥	¥	V	V	¥	¥	∢	¥
PDB TD	lubd	2gli	2gli	2gli	2gli	2gli	2gli	2gli	2gli	2gli
SEQ NO:	485	485	485	485	485	485	485	485	485	485

PDB annotation	ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	RNA BINDING PROTEIN G-PROTEIN, BETA-BARREL	RNA BINDING PROTEIN G-PROTEIN, BETA-BARREL	TRANSLATIONAL GTPASE EF-G RIBOSOMAL TRANSLOCASE, TRANSLATIONAL GTPASE							MYOSIN MYOSIN MOTOR				MYOSIN MYOSIN MOTOR					MUSCLE PROTEIN MDE; MUSCLE	PROTEIN	MUSCLE PROTEIN MDE; MUSCLE
Coumpound	Ď;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ELONGATION FACTOR TU (EF-TU); CHAIN: A, B, C, D	ELONGATION FACTOR TU (EF-TU); CHAIN: A, B, C, D	ELONGATION FACTOR G; CHAIN: NULL;	ACYL-COENZYME A	BINDING PROTEIN ACYL-	COENZYME A BINDING	PROTEIN (ACBP) COMPLEX	COENZYME A (NMR. 20	STRUCTURES) IACA 4	MYOSIN HEAVY CHAIN;	CHAIN: A; MYOSIN REGIII ATORY I IGHT	CHAIN; CHAIN: Y; MYOSIN	ESSENTIAL LIGHT CHAIN; CHAIN: 2:	MYOSIN HEAVY CHAIN;	CHAIN: A; MYOSIN	REGULATORY LIGHT	CHAIN; CHAIN: Y; MYOSIN	ESSENTIAL LIGHT CHAIN; CHAIN: 7:	MYOSIN; CHAIN: A, B, C, D,	E, F, G, H;	MYOSIN; CHAIN: A, B, C, D,
SeqFold score				627.43									510.37									524.14		
PMF		0.94	0.95		-	0.19	0.11	•									-							-
Verify score		0.23	0.03		1.12	-0.05	-0.46	?			-						0.41							0.58
PSI- BLAST		1.40E-34	3.40E-33	0	0	3.40E-17	0.0014						0				0					0		0
End		159	619	454	454	192	1954						794				807					292		66
Start		523	551	58	28	49	1926						-				5					-		2
Chain		⋖	A	٧	A						•		¥	. 			4		_			A	ļ	٨
PDB U		2gli	2gli	1d2e	1d2e	1dar	laca						167t				1671					191		
SEQ NO.		485	485	486	486	486	487				_		487				487					487	į	148/

SEQ SEQ	PDB UD	Chain ID	Start	End	PSI. BLAST	Verify score	PMF score	SeqFold score	Coumpound	PDB annotation
									E, F, G, H;	PROTEIN
487	1br2	¥	11	725	0	0.58			MYOSIN; CHAIN: A, B, C, D, E, F;	MUSCLE PROTEIN MUSCLE PROTEIN
487	1br2	¥	11	739	0			470.06	MYOSIN; CHAIN: A, B, C, D, E, F;	MUSCLE PROTEIN MUSCLE PROTEIN
487	1btk	< −	1394	1495	9.80E-08	-0.2	0.11		BRUTON'S TYROSINE KINASE; CHAIN: A, B;	TRANSFERASE BRUTON'S AGAMMAGLOBULINEMIA TYROSINE KINASE, BTK, TRANSFERASE, PH DOMAIN, BTK MOTIF, ZINC BINDING, X-LINKED 2 AGAMMAGLOBULINEMIA, TYROSINE-PROTEIN KINASE
487	1btn		1216	1306	8.40E-13	0.42	0.28		BETA-SPECTRIN; 18TN 4 CHAIN: NULL; 18TN 5	SIGNAL TRANSDUCTION PROTEIN
487	lbtn		1318	1382	0.00042	-0.39	0.07		BETA-SPECTRIN; 1BTN 4 CHAIN: NULL; 1BTN 5	SIGNAL TRANSDUCTION PROTEIN
487	loii		796	933	2.80E-19	0.21	-0.19		COLICIN IA; CHAIN: NULL;	TRANSMEMBRANE PROTEIN COLICIN, BACTERIOCIN, ION CHANNEL FORMATION, TRANSMEMBRANE 2 PROTEIN
487	1cii		810	616	2.80E-18	0.03	-0.19	,	COLICIN IA; CHAIN: NULL;	TRANSMEMBRANE PROTEIN COLICIN, BACTERIOCIN, ION CHANNEL FORMATION, TRANSMEMBRANE 2 PROTEIN
487	lcun	¥	726	940	1.40E-13	0.07	-0.14		ALPHA SPECTRIN; CHAIN: A, B, C;	STRUCTURAL PROTEIN TWO REPEATS OF SPECTRIN, ALPHA HELICAL LINKER REGION, 2.2 TANDEM 3-HELIX COLLED-COILS, STRUCTURAL PROTEIN
487	Icun	Ą	809	946	1.10E-14	0.27	-0.17		ALPHA SPECTRIN; CHAIN: A, B, C;	STRUCTURAL PROTEIN TWO REPEATS OF SPECTRIN, ALPHA HELICAL LINKER REGION, 2.2 TANDEM 3-HELIX COILED-COILS, STRUCTURAL PROTEIN
487	¥FP1	¥	3	807	0	0.23	-		MYOSIN HEAD; CHAIN: A; MYOSIN HEAD; CHAIN: Y; MYOSIN HEAD; CHAIN: Z;	CONTRACTILE PROTEIN MYOSIN MOTOR, CONFORMATIONAL CHANGES
487	lefi	V	1710	2002	4.20E-27	0.13	66.0		MOESIN; CHAIN: A, B; MOESIN; CHAIN: C, D;	MEMBRANE PROTEIN CRYSTAL STRUCTURE, MEMBRANE, FERM DOMAIN, TAIL DOMAIN
487	lez3	Ą	809	885	1.40E-11	0.37	-0.2		SYNTAXIN-1A; CHAIN: A. B,	ENDOCYTOSIS/EXOCYTOSIS

			-4		<u> </u>			1
PDB annotation	SYNAPTOTAGMIN ASSOCIATED 35 KDA PROTEIN, P35A, THREE HELIX BUNDLE	ENDOCYTOSIS/EXOCYTOSIS SYNAPTOTAGMIN ASSOCIATED 35 KDA PROTEIN, P35A, THREE HELIX BUNDLE	SIGNALING PROTEIN DAPP1, PHISH, BAM32; PLECKSTRUN, 3-PHOSPHONOSITIDES, INOSITOL TETRAKISPHOSPHATE 2 SIGNAL TRANSDUCTION PROTEIN, ADAPTOR PROTEIN	SIGNALING PROTEIN DAPP1, PHISH, BAM32; PLECKSTRIN, 3-PHOSPHOINOSTIDES, INOSITOL TETRAKISPHOSPHATE 2 SIGNAL TRANSDUCTION PROTEIN, ADAPTOR PROTEIN	SIGNALING PROTEIN DAPPI, PHISH, BAM32; PLECKSTRIN, 3-PHOSPHONOSITIDES, INOSITOL TETRAKISPHOSPHATE 2 SIGNAL TRANSDUCTION PROTEIN, ADAPTOR PROTEIN	SIGNALING PROTEIN DAPP1, PHISH, BAM32; PLECKSTRIN, 3-PHOSPHOINOSITIDES, INOSITOL TETRAKISPHOSPHATE 2 SIGNAL TRANSDUCTION PROTEIN, ADAPTOR PROTEIN, ADAPTOR PROTEIN	SIGNALING PROTEIN DAPPI, PHISH, BAM32; PLECKSTRIN, 3-PHOSPHOINOSITIDES, INOSITOL TETRAKISPHOSPHATE 2 SIGNAL TRANSDUCTION PROTEIN, ADAPTOR PROTEIN	SIGNALING PROTEIN DAPPI, PHISH, BAM32; PLECKSTRIN, 3-PHOSPHOINOSITIDES, INOSITOL TETRAKISPHOSPHATE 2 SIGNAL
Coumpound	;o	SYNTAXIN-1A; CHAIN: A, B, C,	DUAL ADAPTOR OF PHOSPHOTYROSINE AND 3- CHAIN: A;	DUAL ADAPTOR OF PHOSPHOTYROSINE AND 3- CHAIN: A;	DUAL ADAPTOR OF PHOSPHOTYROSINE AND 3- CHAIN: A;	DUAL ADAPTOR OF PHOSPHOTYROSINE AND 3- CHAIN: A;	DUAL ADAPTOR OF PHOSPHOTYROSINE AND 3 CHAIN: A;	DUAL ADAPTOR OF PHOSPHOTYROSINE AND 3- CHAIN: A;
SeqFold score				·				
PMF score		-0.18	0.92	96.0	0.36	0.99	98.0	0.25
Verify score		0.3	0.43	0.13	0.19	0.54	-0.06	-0.03
PSI- BLAST		1.10E-16	2.80E-22	7.00E-10	4.20E-10	5.60E-22	8.40E-10	1.40E-10
End AA		933	1308	1382	1497	1308	1382	1497
Start		814	1215	1331	1397	1215	1331	1397
Chain U		<	⋖ .	∢	∀	Ą.	A	∢
PDB		lez3	Ifao	Ifao	lfao	1fb8	1tb8	1408
SEQ No.	·	487	487	487	487	487	487	487

PDB annotation	TRANSDUCTION PROTEIN, ADAPTOR PROTEIN	SIGNALING PROTEIN ARFI GUANINE NUCLEOTIDE ÉXCHANGE FACTOR AND PH DOMAIN	CELL ADHESION 3 SUBDOMAINS,CYTOSKELETON, CELL ADHESION	CONTRACTILE PROTEIN MYOSIN, DICTYOSTELIUM, MOTOR, MANT, ATPASE, ACTIN-BINDING, 2 COILED COIL	CONTRACTILE PROTEIN MYOSIN, DICTYOSTELIUM, MOTOR, MANT, ATPASE, ACTIN-BINDING, 2 COILED COIL	CONTRACTILE PROTEIN ATPASE, MYOSIN, COLLED COIL, ACTIN- BINDING, ATP-BINDING, 2 HEPTAD REPEAT PATTERN, METHYLATION, ALKYLATION, 3 PHOSPHORYLATION, CONTRACTILE PROTEIN	CONTRACTILE PROTEIN ATPASE, MYOSIN, COILED COIL, ACTIN- BINDING, ATP-BINDING, 2 HEPTAD REPEAT PATTERN, METHYLATION, ALKYLATION, 3 PHOSPHORYLATION, CONTRACTILE PROTEIN	·	,
Coumpound		GRP1; CHAIN: A;	RADIXIN; CHAIN: A;	MYOSIN; CHAIN: NULL;	MYOSIN; CHAIN: NULL;	MYOSIN; CHAIN: NULL;	MYOSIN; CHAIN: NULL;	PHOSPHORYLATION PLECKSTRIN (N-TERMINAL PLECKSTRIN HOMOLOGY DOMAIN) MUTANT IPLS 3 WITH LEU GLU (HIS)6 ADDED TO THE C TERMINUS IPLS 4 (INS(G105-LEHHHHHHH)) (NMR, 25 STRUCTURES) IPLS 5	PHOSPHORYLATION PLECKSTRIN (N-TERMINAL PLECKSTRIN HOMOLOGY DOMAIN) MUTANT IPLS 3
SeqFold score				540.6		479.56			
PMF score		0.31	 	_	-			0.46	0.04
Verify score		0.21	0.08	·	0.6	·	0.63	0.11	-0.55
PSI- BLAST		2.80E-16	2.80E-26	0	0	0	0	2.80E-17	5.60E-05
End		1308	2044	739	725	671	671	1315	1381
Start		1215	1710	2	5	-	s	1214	1342
Chain ID		∢	V				٠		
PDB ID		1fgy	1gc7	1lvk	11× 1×	d d	1mi d	Ipis	1pls
SEQ B G S		487	487	487	487	487	487	487	487

PDB annotation			SIGNAL TRANSDUCTION SON OF SEVENLESS, PLECKSTRIN, SON OF SEVENLESS, SIGNAL TRANSDUCTION	SIGNAL TRANSDUCTION SON OF SEVENLESS; PLECKSTRIN, SON OF SEVENLESS, SIGNAL TRANSDUCTION	SIGNAL TRANSDUCTION IRS-1; BETA-SANDWHICH, SIGNAL TRANSDUCTION	CONTRACTILE PROTEIN TRIPLE- HELIX COILED COIL, CONTRACTILE PROTEIN	MUSCLE PROTEIN MUSCLE PROTEIN, MYOSIN SUBFRAGMENT-1, MYOSIN HEAD, 2 MOTOR PROTEIN	MUSCLE PROTEIN MUSCLE PROTEIN, MYOSIN SUBFRAGMENT-1, MYOSIN HEAD, 2 MOTOR PROTEIN	MICROTUBULES MICROTUBULES, ALPHA-TUBULIN, BETA-TUBULIN, GTPASE HELIX	MICROTUBULES MICROTUBULES, ALPHA-TUBULIN, BETA-TUBULIN, GTPASE HELIX	KINASE KINASE, SIGNAL
Coumpound	WITH LEU GLU (HIS)6 ADDED TO THE C TERMINUS 1PLS 4 (INS(G10S-LEHHHHHH)) (NMR, 25 STRUCTURES) 1PLS 5	PHOSPHORYLATION PLECKSTRIN (N-TERMINAL PLECKSTRIN (N-TERMINAL PLECKSTRIN HOMOLOGY DOMAIN) MUTANT IPLS 3 WITH LEU GLU (HIS)6 ADDED TO THE C TERMINUS IPLS 4 (INS(G105-LEHHRICH)) (NMR, 25 STRUCTURES) IPLS 5	SOS 1; CHAIN: NULL;	SOS 1; CHAIN: NULL;	NSULIN RECEPTOR SUBSTRATE 1: CHAIN: A, B;	HUMAN SKELETAL MUSCLE ALPHA-ACTININ 2; CHAIN: A;	MYOSIN; CHAIN: A, B, C;	MYOSIN; CHAIN: A, B, C;	TUBULIN; CHAIN: A, B;	TUBULIN; CHAIN: A, B;	CALCIUM/CALMODULIN-
SeqFold score							419.33		727.18		
PMF score		-0.01	-0.01	0.05	0.25	-0.13		1		_	
Verify		0.02	0.01	-0.14	-0.27	0.05		0.53		0.8	0.35
PSI- BLAST		9.80E-12	8.40E-15	0.00014	7.00E-07	1.30E-20	0	0	0	0	1.50E-87
End		1495	1308	1381	1464	973	801	277	440	440	312
Start		1394	1211	1331	1331	797	2	4	_	-	16
Chain ID					∀	٧	4	Ą	V	V V	
PDB UD		1pls	1 pms	Ipms	1998	lquu	2mys	2mys	Itub	1tub	1a06
SEQ SO SO SO SO SO SO SO SO SO SO SO SO SO		487	487	487	487	487	487	487	489	489	492

	_				
PDB annotation	TRANSDUCTION, CALCIUM/CALMODULIN	KINASE KINASE, SIGNAL TRANSDUCTION, CALCIUM/CALMODILI.N	TRANSFERASE TRANSFERASE, SERINE/THREONINE-PROTEIN KINASE, CASEIN KINASE, 2 SER/THR KINASE		PROTEIN KINASE CDK2; PROTEIN KINASE, CELL CYCLE, PHOSPHORYLATION, STAUROSPORINE, 2 CELL DIVISION,
Coumpound	DEPENDENT PROTEIN KINASE; CHAIN: NULL;	CALCIUM/CALMODULIN- DEPENDENT PROTEIN KINASE: CHAIN: NULL:	PROTEIN KINASE CK2/ALPHA-SUBUNIT; CHAIN: NULL;	TRANSFERASE(PHOSPHOTR ANSFERASE) \$C-AMP\$- DEPENDENT PROTEIN KNASE (E.C.7.11.37) (\$CAPK\$) 1APM 3 (\$CAPK\$) 1APM 3 (CATALYTIC SUBUNIT) ALPHA ISOENZYME MUTANT WITH SER 139 1APM 4 REPLACED BY ALA (\$(\$1394\$), COMPLEX WITH THE PEPTINE 1APM 5 INHIBITOR PKI(5-24) AND THE DETERGENT MEGA-8 IAPM 6 IAPM 6 IAPM 6 IAPM 5 IAPM 6 IAPM 6 IAPM 5 IAPM 6 IAPM 5 IAPM 6 IAPM 5 IAPM 6 IAPM 7 IAPM 1SOENZYME MUTANT WITH SER 139 IAPM 4 ICALALYTIC SUBUNIT) ALPHA 1SOENZYME MUTANT WITH SER 139 IAPM 4 REPLACED BY ALA (\$(\$1394\$), COMPLEX WITH THE PEPTIDE 1APM 5 INHIBITOR PKI(5-24) AND THE DETERGENT MEGA-8 INHIBITOR PKI(5-24) AND	CYCLIN-DEPENDENT PROTEIN KINASE 2; CHAIN: NULL;
SeqFold		121.63	90.87	158.35	120.73
PMF	ж.			-	
Verify score				0.41	
PSI- BLAST		1.50E-87	1.70E-43	0 0	1.00E-57
End AA		318	316	333	314
Start AA		17	2		22
Chain ID				рд рд	
PDB ID		1a06	Ia6o	n n lap	laq1
SEQ ID NO:		492	492	492	492

PDB annotation	MITOSIS, INHIBITION	COMPLEX (KINASE/INHIBITOR) CDK6; P19INK4D; CYCLIN DEPENDENT KINASE, CYCLIN DEPENDENT KINASE, INHIBITORY 2	PROTEIN, CDK, INK4, CELL CYCLE, COMPLEX (KINASE/INHIBITOR) HEADER HELIX	COMPLEX (INHIBITOR PROTEIN/KINASE) INHIBITOR PROTEIN, CYCLIN-DEPENDENT	KINASE, CELL CYCLE 2 CONTROL, ALPHA/BETA, COMPLEX (INHIBITOR PROTEIN/KINASE)	TRANSFERASE CSK; PROTEIN KINASE C-TERMINAL SRCKINASE	PHOSPHORYLATION, 2 STAUROSPORINE, TRANSFERASE	PHOSPHOTRANSFERASE PROTEIN KINASE 1CKI 18	=						,								
Coumpound		CYCLIN-DEPENDENT KINASE 6; CHAIN: A, C; CYCLIN-DEPENDENT KINASE INHIBITOR; CHAIN:	B, D;	CYCLIN-DEPENDENT KINASE 6; CHAIN: A; P19INK4D; CHAIN: B;		C-TERMINAL SRC KINASE;	(T	CASEIN KINASE I DELTA; ICKI 6 CHAIN: A. B. ICKI 7	PHOSPHOTRANSFERASE	CAMP-DEPENDENT	CATAL YTIC SUBUNIT ICMK	3 (E.C.2.7.1.37) ICMK 4	PHOSPHOTRANSFERASE CAMP-DEPENDENT	PROTEIN KINASE	3 (E.C.2.7.1.37) ICMK 4	TRANSFERASE(PHOSPHOTR	DEPENDENT PROTEIN	KINASE (E.C.2.7.1.37) (CAPK)	1CTP 3 (CATALYTIC SUBUNIT) ICTP 4	TRANSFERASE(PHOSPHOTR	ANSFERASE) CAMP-	KINASE (E.C.2.7.1.37) (CAPK)	1CTP 3 (CATALYTIC
SeqFold score		117.12		139.32		120.46		82.9					156.27	•						152.26			
PMF																1						0	
Verify							•		0.42		•					0.32							
PSI- BLAST		2.80E-54		1.40E-59		1.40E-39		9.80E-51	0				0	•		0				0			
End		303	•	308		286		303	315				333			315				330			
Start		23		18		18		17	_				m ,			-				3			_]
Chain ID		4		4		¥		V	E				ш			Ξ				Э			
PDB		1bi8		XI4I		1byg		1cki	1cm	<u>~</u>			두 <u>무</u>			1ctp				1ctp		_	
SEQ NO:		492		492		492		492	492				492			492				492			

	_			, 		т		·
PDB annotation		TRANSFERASE KINASE DOMAIN, AUTOINHIBITORY FRAGMENT, HOMODIMER	TRANSFERASE KINASE DOMAIN, AUTOINHIBITORY FRAGMENT, HOMODIMER	PHOSPHOTRANSFERASE FGFRIK, FIBROBLAST GROWTH FACTOR RECEPTOR 1; TRANSFERASE, TYROSINE-PROTEIN KINASE, ATP- BINDING, 2 PHOSPHORYLATION, RECEPTOR, PHOSPHOTRANSFERASE	PHOSPHOTRANSFERASE FGFRIK, FIBROBLAST GROWTH FACTOR RECEPTOR 1: TRANSPERASE, TYROSINE-PROTEIN KINASE, ATP- BINDING, 2 PHOSPHORYLATION, RECEPTOR, PHOSPHOTRANSFERASE	PROTEIN KINASE CDK2; TRANSFERASE, SERINE/THREONINE PROTEIN KINASE, ATP-BINDING, 2 CELL CYCLE, CELL DIVISION, MITOSIS, PHOSPHORYLATION	SERINE/THREONINE-PROTEIN KINASE CSBP, RK, P38; PROTEIN SER/THR-KINASE, SERNJETHREONINE-PROTEIN KINASE	COMPLEX (TRANSFERASE/SUBSTRATE) TYROSINE KINASE, SIGNAL TRANSDUCTION, PHOSPHOTRANSFERASE, 2 COMPLEX (KINASEPPEPTIDE SUBSTRATEATP
Coumpound	SUBUNIT) 1CTP 4	SERINE/THREONINE- PROTEIN KINASE PAK- ALPHA; CHAIN! A, B; SERINE/THREONINE- PROTEIN KINASE PAK- ALPHA; CHAIN! C, D;	SERNETHREONINE- PROTEIN KINASE PAK- ALPHA; CHAIN! A, B; SERINETHREONINE- PROTEIN KINASE PAK- ALPHA; CHAIN! C, D;	FGF RECEPTOR 1; CHAIN: A, B;	FGF RECEPTOR I; CHAIN: A, B;	HUMAN CYCLIN- DEPENDENT KINASE 2; CHAIN: NULL;	P38 MAP KINÁSE; CHAIN: NULL;	INSULIN RECEPTOR; CHAIN: A; PEPTIDE SUBSTRATE; CHAIN: B;
SeqFold score				123.75	127.9	141.29	104.86	105.26
PMF			-					
Verify score		0.38	0.21		24			
PSI- BLAST		2.80E-69	5.10E-69	2.80E-38	1.20E-40	1.40E-60	2.80E-45	9.80E-40
End		303	293	286	285	314	346	297
Start		23	4		12	22	s	6
Chain D		U .	U	∢	m			<
PDB		113m	113m	1fgk	lfgk	1hcl	lian	Lir3
SEQ NO ID		492	492	492	492	492	492	492

												
PDB annotation	ANALOG), ENZYME, 3 COMPLEX (TRANSFERASE/SUBSTRATE)	TRANSFERASE INK3; TRANSFERASE, INK3 MAP KINASE, SERINE/THREONINE PROTEIN 2 KINASE	KINASE KINASE, TWITCHIN, INTRASTERIC REGULATION	KINASE KINASE, TWITCHIN, INTRASTERIC REGULATION	KINASE KINASE, TWITCHIN, INTRASTERIC REGULATION	TRANSFERASE MITOGEN ACTIVATED PROTEIN KINASE; SERNETHREONINE-PROTEIN KINASE, 2 P38	KINASE RABBIT MUSCLE PHOSPHORYLASE KINASE; GLYCOGEN METABOLISM, TRANSFERASE, SERNE/THREONINE-PROTEIN, 2 KINASE, ATP-BINDING, CALMODULIN-BINDING,	KINASE RABBIT MUSCLE PHOSPHORYLASE KINASE; GLYCOGEN METABOLISM, TRANSFERASE, SERNE/THREONINE- PROTEIN. 2 KINASE, ATP-BINDING, CALMODULIN-BINDING	TRANSFERASE MAP KINASE, SERINE/THREONINE PROTEIN KINASE, TRANSFERASE	SERINE KINASE SERINE KINASE, TITIN, MUSCLE, AUTOINHIBITION	SERINE KINASE SERINE KINASE, TITIN, MUSCLE, AUTOINHIBITION	TRANSFERASE MITOGEN ACITVATED PROTEIN KINASE, MAP 2, ERK2; TRANSFERASE, SERINETHREONINE-PROTEIN KINASE, MAP KINASE, 2 ERK2
Coumpound		C-JUN N-TERMINAL KINASE; CHAIN: NULL;	TWITCHIN; CHAIN: NULL;	TWITCHIN; CHAIN: A, B;	TWITCHIN; CHAIN: A, B,	MAP KINASB P38; CHAIN: NULL;	PHOSPHORYLASE KINASE; CHAIN: NULL;	PHOSPHORYLASE KINASE; CHAIN: NULL;	ERKZ; CHAIN: NULL;	TITIN; CHAIN: A, B;	TITIN; CHAIN: A, B;	EXTRACELLULAR REGULATED KINASE 2; CHAIN: NULL;
SeqFold score		127.36		139.94		121.52	109.51		118.65	126.69		130.2
PMF					-			_			. 1	
Verify score			0.27		0.4			9.6			0.49	
PSI- BLAST		7.00E-54	1.70E-70	3.40E-71	3.40E-71	1.40E-56	1.20E-81	1.20E-81	5.60E-50	4.20E-65	4.20E-65	4.20E-56
End		357	334	351	284	350	282	279	341	334	274	346
Start AA		∞	22		17	4	22	23	18	19	22	11
Chain B B				¥	۷.					٧	< │	
PDB D		Įi,	lkoa	1kob	1kob	1p38	lphk	1 phk	lpm c	1tki	igi i	3erk
SEQ No B		492	492	492	492	492	492	492	492	492	492	492

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PDB annotation	LIPID TRANSPORT APO A-1; LIPOPROTEIN, LIPID TRANSPORT, CHOLESTEROL METABOLISM, 2 ATHEROSCLEROSIS, HDL, LCAT- ACTIVATION	TRANSFERASE IL-2-INDUCIBLE T- CELL KINASE; TRANSFERASE, REGULATORY INTRAMOLECULAR COMPLEX, KINASE	COMPLEX (ADAPTOR PROTEIN/PEPTIDE) ASH, GROWTH PACTOR RECEPTOR-BOUND PROTEIN 2; COMPLEX (ADAPTOR PROTEIN/PEPTIDE), SH3 DOMAIN, 2 GUANINE-NUCLEOTIDE RELEASING FACTOR	COMPLEX (TRANSFERASE/PEPTIDE) COMPLEX (TRANSFERASE/PEPTIDE), SIGNAL TRANSDUCTION, 2 SH3 DOMAIN			ENDOCYTOSIS/EXOCYTOSIS SYNAPTOTAGMIN ASSOCIATED 35 KDA PROTEIN, P35A, THREE HELIX BUNDLE	MEMBRANE PROTEIN FOUR HELIX BUNDLE, ALPHA HELIX	TRANSFERASE PROTO-ONCOGENE TYROSINE KINASE; PROTO- ONCOGENE, TRANSFERASE,
Commbonuq	APOLIPOPROTEIN A-I; CHAIN: A, B, C, D;	ITK; CHAIN: NULL;	GRB2; CHAIN: A; SOS; CHAIN: B;	ABL TYROSINE KINASE; CHAIN: A, C, E, G; PEPTIDE P41; CHAIN: B, D, F, H;	COMPLEX (ONCOGENE PROTEIN/PEPTIDE) C-CRK (N-TERMINAL SH3 DOMAIN) (C-CRKSH3-N) COMPLEXED WITH ICKA 3 GG PEPTIDE (PRO-PRO-PRO-ALA-LEU- PRO-PRO-PSO-ALA-LEU- PRO-PRO-LYS-LYS-ARG)	PHOSPHOTRANSFERASE C- SKC KINASE (SH3 DOMAIN) (E.C.2.7.1.112) 1CSK 3	SYNTAXIN-1A; CHAIN: A, B, C;	SSOI PROTEIN; CHAIN: A;	PHOSPHOTRANSFERASE FYN; CHAIN: A; 3BP-2; CHAIN: B;
SeqFold score	66.14				,				
PMF score	7	0.75	П	0.95	66.0	0.99	0.05	0.05	0.88
Verify score		0.64	0.47	0.67	0.58	0.72	0.22	0.24	0.02
PSI- BLAST	0.0042	1.70E-08	4.20E-19	7.00E-18	8.40E-18	9.80E-19	2.80E-06	0.0056	2.80E-17
End AA	278	440	442	444	44.	442	214	256	444
Start AA	58	370	388	390	389	387	93	166	385
Chain ID	Ą		4	4	⋖	∢	<	4	A
PDB CD	laví	lawj	laze	1bbz	1 cka	1csk	lez3	1fio	1fyn
SEQ No:	493	493	493	493	493	493	493	493	493

PDB annotation	FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGERDNA) COMPLEX (ZINC FINGERDNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA FINGER, PROTEIN DESIGN, 2 CRYSTAL, STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION. PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CEYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CXYSTAL STRUCTURE, COMPLEX ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CXYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC
Coumpound	OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE, CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E;
SeqFold score								110.98	,	
PMF			0.77	-	<u>-</u>	-	-		_	0.87
Verify score		0.03	0.12	0.38	0.4	0.24	0.34		0.05	-0.05
PSI- BLAST		5.60E-35	1.40E-33	6.80E-49	1.70E-50	3.40E-51	5.10E-51	3.40E-51	3.40E-47	4.20E-36
End AA		478	505	338	366	394	422	423	477	505
Start AA		370	398	257	285	313	341	341	397	397
Chain ID		V	V	U	U	U	U	ပ	ប	၁
PDB		lalh	laih	J me	y y	J me	Jme y	Ime y	Ime y	1mc
SEQ NO B		498	498	498	498	498	498	498	. 498	498

								
PDB annotation	FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC · FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INITIATION ZNIC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INITATION ZINC FINGER PROTEN	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA
Coumpound	CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	TFIIIA; CHAIN: A, D, 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	TFIIIA; CHAIN: A, D, 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	TFIIIA; CHAM: A, D; 5S RIBOSOMAL RNA GENE; CHAM: B, C, E, F;
SeqFold score	·					102.44		
PMF score		-	_	-0.15	-0.01		0.94	0.92
Verify score		0.23	0.19	0.1	0.00		0.11	-0.02
PSI- BLAST		1.00臣-49	1.70E-35	3.40E-12	1.00E-33	1.40E-68	1.70E-36	8.50E-39
End		505	511	282	375	447	437	487
Start		424	452	255	221	285	286	342
Chain ID		ပ	ပ	ប	⋖	⋖	∢	¥ .
PDB CI	>	1me y	Jme y	Ime y	1476	146	1476	146
SEQ S		498	498	498	498	498	498	498

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PDB annotation	POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION	REGULATION/DNA), RNA	POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION	REGULATION/DNA) YING-YANG 1;	INTIATOR ELEMENT, YYI, ZINC 2	FINGER PROTEIN, DNA-PROTEIN	RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGIII ATTOMONA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA) YING-YANG 1;	TRANSCRIPTION INITIATION,	INITIATOR ELEMENT, YY1, ZINC 2	RECOGNITION 3 COMPLEY	(TRANSCRIPTION REGULATION DNA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA) YING-YANG 1;	TRANSCRIPTION INITIATION,	INITIATOR ELEMENT, YYI, ZINC 2	PECOGNITION 2 COMBINEY	(TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION	TEGULATION/DNA) YING-YANG I;	INTITATOR ELEMENT YYL ZINC?	FINGER PROTEIN, DNA-PROTEIN	RECOGNITION, 3 COMPLEX	(TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA) YING-YANG 1;	INANSCRIPTION INITIATION,	FINGER PROTEIN, DNA-PROTEIN
Coumpound		TFIIIA; CHAIN: A, D; SS RIBOSOMAL RNA GENE; CHAIN: B, C, F, F.	יין אין אין אין אין אין אין אין אין אין		YY1; CHAIN: C; ADENO-	ASSOCIATED VIRUS PS	CHAIN: A, B;			YY1; CHAIN: C; ADENO-	ASSOCIATED VIRUS P5	INITIATOR ELEMENT DNA;	CHAIN: A, B;			YY1; CHAIN: C; ADENO-	ASSOCIATED VIRUS PS	INITIATOR ELEMENT DNA,	CHAIN: A, B;			YY1; CHAIN: C; ADENO-	ASSOCIATED VIKUS PS	CHAIN: A. B.				YY1; CHAIN: C; ADENO-	ASSOCIATED VIRUS PS	CHAIN: A B.	(7 ty 17)
SeqFold																88.23															
PMF		0.95			0.05					_									_			1						_			
Verify score		0.14			-0.02					0.25												0.23						0.36			
PSI- BLAST		8.50E : 35			1.00E-30		•			1.70E-34						2.80E-56				•		2.80E-56		•				8.50E-35			
End AA		205	•		338					366						395						394					1	394			
Start AA		370	_		228					265						285						290						567			
Chain ID		4			ပ					ပ						ပ						ပ						ر			
PDB ID		9,111			Jubd					1ubd			.,			PqnI					+	Pani		_			┰	pon			
SEQ NO:		498			498					498						498		_	•			498					8	984			

Coumpound PDB annotation	RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	YYI; CHAIN: C; ADENO- COMPLEX (TRANSCRIPTION ASSOCIATED VIRUS P5 REGULATION/DNA) YING-YANG 1; ATAINSCRIPTION INTIATION, CHAIN-R ELEMENT YYI, ZINC2	YA;	INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN PECOCENTION 3 COMPIEY	(TRANSCRIPTION REGULATION/DNA)	ZINC FINGER PROTEIN GLII; COMPLEX (DNA-BINDING CHAIN: C, PROTEIN NA) FIVE-FINGER GLI; GLI, PROTEIN CON THE CONTROL OF THE CONT		11;	D; CHAIN: A; DNA; CHAIN: C, ZINC FINGER, COMPLEX (DNA-	BINDING PROTEIN/DNA)	<u>:</u>	CHAIN: A; DNA; CHAIN: C, PROTEIN/DNA) FIVE-FINGER GLI; GLI, D: ZINC FINGER, COMPLEX (DNA-	BINDING PROTEIN/DNA)	:11		CHAIN: A; DNA; CHAIN: C, PROTEIN/DNA) FIVE-FINGER GLI; GLI,		=					
Coum		YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DI CHAIN: A B:	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DI	CHAIN: A, B;		ZINC FINGER CHAIN: A; DN	.; 	ZINC FINGER	CHAUN: A; DN.		ZINC FINGER	CHAIN: A; DN D:		ZINC FINGER	CHAIN: A; DN		ú	D; ZINC FINGER	D; ZINC FINGER CHAIN: A; DN	D; ZINC FINGER CHAIN: A; DN D;	D; ZINC FINGER CHAIN: A; DN. D; ZINC FINGER	D; ZNC FINGER CHAIN: A; DN. D; ZINC FINGER CHAIN: A; DN.	D; ZINC FINGER CHAIN: A; DN. D; ZINC FINGER CHAIN: A; DN. D;
SeqFold score								97.78	_				_										
PMF		-	 0.92			0.28					_			_				0.86	98.0	0.86	0.86	0.86	0.86
Verify score		0	-0.07			-0.26					0.24			0.36				0.14	0.14	0.14	0.14	0.14	0.14
PSI- BLAST		2.80E-56	 6.80E-36			8.50E-33		2.80E-70			2.80E-70			3.40E-33				1.30E-63	1.30E-63	1.30E-63	1.30E-63	1.30E-63 4.20E-61	1.30E-63 4.20E-61
End AA		422	 202			337		423			422			423				479	479	479	479	507	507
Start		311	 405			200		285			290			293				313	313	313	313	313	313
Chain ID		O	 ပ			4		٧			A			٧				∢	4	∀	4 4	4 4	4 4
PDB CD		Iubd	lubd			2gli		2gli			2gli			2gli				2gli	2gli	2gli	2gli 2gli	2gli 2gli	2gli 2gli
SEQ EQ		498	498			498		498			498			498				498	498	498	498	498	498

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PDB annotation	PROTEIN/DNA) FIVE-FINGER GLI, GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	LIGASE CYCLIN A/CDK2- ASSOCIATED PROTEIN P45; CYCLIN A/CDK2-ASSOCIATED PROTEIN P19; SKP1, SKP2, F-BOX, LRK, LEUCINE- RICH REPEAT, SCF, UBIQUITIN, 2 E3, UBIQUITIN PROTEIN LIGASE	LIGASE SKP2 F-BOX; SKP1; SKP1, SKP2, F-BOX, LRR, LEUCINE-RICH REPEAT, SCF, UBIQUITIN, 2 E3, UBIQUITIN PROTEIN LIGASE		LIGASE CBL, UBCH7, ZAP-70, E2, UBIQUITIN, E3, PHOSPHORYLATION, 2 TYROSINE KINASE, UBIQUITINATION, PROTEIN DEGRADATION,	TRANSCRIPTION REGULATION PROTO-ONCOGENE, NUCLEAR BODIES (PODS), LEUKEMIA, 2 TRANSCRIPTION REGULATION			LIGASE CBL, UBCH7, ZAP-70, E2,	UBIQUITIN, E3, PHOSPHORYLATION, 2 TYROSINE KINASE,	UBIQUITINATION, PROTEIN DEGRADATION.	
Coumpound	CHAIN: A; DNA; CHAIN: C, D;	SKP2; CHAIN: A, C, E, G, I, K, M, O; SKP1; CHAIN: B, D, F, H, J, L, N, P;	CYCLIN A/CDK2. ASSOCIATED P19; CHAIN: A, C; CYCLIN A/CDK2. ASSOCIATED P45; CHAIN: B, D;		SIGNAL TRANSDUCTION PROTEIN CBL; CHAIN: A; 2AP-70 PEPTIDE, CHAIN: B; UBIQUITIN-CONJUGATING ENZYME E12-18 KDA UBCH7; CHAIN: C;	TRANSCRIPTION FACTOR PML; CHAIN: NULL;	VIRUS EQUINE HERPES VIRUS-1 (C3HC4, OR RING DOMAIN) ICHC 3 (NMR, 1 STRUCTURE) ICHC 4	VIRUS EQUINE HERPES VIRUS-1 (C3HC4, OR RING DOMAIN) 1CHC 3 (NMR, 1 STRUCTURE) 1CHC 4	SIGNAL TRANSDUCTION	PROTEIN CBL; CHAIN: A; ZAP-70 PEPTIDE; CHAIN: B;	UBIQUITIN-CONJUGATING ENZYME E12-18 KDA	UBCH7; CHAIN: C;
SeqFold score			·									
PMF		0.52	0.53		0.16	0.01	0.53	69.0	0.81			
Verify score		-0.78	-0.81		0.08	-0.37	-0.07	0.02	-0.55			
PSI- BLAST		0.0007	2.80E-06		1.70E-06	1.40E-13	8.40E-17	3.40E-16	5.60E-11			
End AA			57		907	305	323	316	307			
Start AA		28	28		998	255	259	261	797			
Chain Et in		4	₹		∢				¥			
PDB CD		Ifqv	15:1		Ifbv	Ibor	1chc	1chc	1fbv			
SEQ EQ		499	499		200	105	501 ·	501	201			

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PDB annotation	LIGASE CBL, UBCH7, ZAP-70, E2, UBIQUITIN, E3, PHOSPHOR YLATION, 2 TYROSINE KINASE, UBIQUITINATION, PROTEIN DEGRADATION,	METAL BINDING PROTEIN RING FINGER PROTEIN MATI; RING FINGER (C3HC4)	DNA-BINDING PROTEIN V(D)J RECOMBINATION ACTIVATING PROTEIN 1; RAG1, V(D)J RECOMBINATION, ANTIBODY, MAD, RING FINGER, 2 ZINC BINUCLEAR CLUSTER, ZINC FINGER, DNA- BINDING PROTEIN	DNA-BINDING PROTEIN V(D)J RECOMBINATION ACTIVATING PROTEIN I; RAGI, V(D)J RECOMBINATION, ANTIBODY, MAD, RING FINGER, 2 ZNC BINUCLEAR CLUSTER, ZINC FINGER, DNA- BINDING PROTEIN		MEMBRANE PROTEIN LECTIN-LIKE, NEUROBIOLOGY, CELL-CELL ADHESION, CELL-CELL 2 RECOGNITION, ALTERNATIVE SPLICING, MEMBRANE PROTEIN	TRANSFERASE DINUCLEOTIDE- BINDING MOTIF, PHOSPHORIBOSYL TRANSFERASE	TRANSFERASE DINUCLEOTIDE- BINDING MOTIF, PHOSPHORIBOSYL TRANSFERASE	TRANSFERASE DINUCLEOTIDE- BINDING MOTIF, PHOSPHORIBOSYL TRANSFERASE	ZINC METALLOPROTEASE P. AERUGINOSA ALKALINE PROTEASE; IKAP 6 CALCIUM BINDING PROTEIN
Coumpound	SIGNAL TRANSDUCTION PROTEIN CBL; CHAIN: A; ZAP-70 PEPTIDE, CHAIN: B; UBIQUITIN-CONJUGATING ENZYME E12-18 KDA UBCH7; CHAIN: C;	CDK-ACTIVATING KINASE ASSEMBLY FACTOR MAT1; CHAIN: A;	RAGI; CHAIN: NULL;	RAGI; CHAIN: NULL;	*	NEUREXIN-I BETA, CHAIN: A, B, C, D, B, F, G, H;	NICOTINATE MONONUCLEOTIDE:5,6- CHAIN: A;	NICOTINATE MONONUCLEOTIDE:5,6- CHAIN: A;	NICOTINÁTE MONONUCLEOTIDE:5,6- CHAIN: A;	ALKALINE PROTEASE; IKAP 4 CHAIN: P; IKAP 5 TETRAPEPTIDE (GLY SER
SeqFold score										
PMF		0.22	0.94	0.45		-0.11	-0.2	-0.18	-0.2	-0.2
Verify score	-0.24	-0.17	-0.11	-0.27		0.21	0.3	0.51	0.56	96.0
PSI- BLAST	1.70E-09	7.00E-13	8.40E-18	1.70E-09		1.40E-17	1.10E-23	1.30E-20	1.10E-23	7.00E-14
End	310	319	333	342		224	1380	1072	1241	1439
Start	263	259	239	263		50	1042	753	116	1076
Chain ED	V	4				A	A	¥	А	Ъ
PDB ID	1fbv	1g25	1 гт	1rmd		lc4r	1d0s	1d0s	1d0s	lkap
SEQ B	501	201	501	105		502	502	502	502	502

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PDB annotation	1KAP 19	ZINC METALLOPROTEASE P. AERUGINOSA ALKALINE PROTEASE; IKAP 6 CALCIUM BINDING PROTEIN IKAP 19	ZINC METALLOPROTEASE P. AERUGINOSA ALKALINE BROTEASE; IKAP 6 CALCIUM BINDING PROTEIN IKAP 19	ZINC METALLOPROTEASE P. AERUGINOSA ALKALINE PROTEASE; IKAP 6 CALCIUM BINDING PROTEIN IKAP 19	OUTER MEMBRANE PROTEIN OSMOPORIN; OUTER MEMBRANE PROTEIN, NON-SPECIFIC PORIN, OSMOPORIN, 2 BETA-BARREL, TRANSMEMBRANE	OUTER MEMBRANE PROTEIN OSMOPORIN; OUTER MEMBRANE PROTEIN, NON-SPECIFIC PORIN, OSMOPORIN, 2 BETA-BARREL, TRANSMEMBRANE	OUTER MEMBRANE PROTEIN OSMOPORIN, OUTER MEMBRANE PROTEIN, NON-SPECIFIC PORIN, OSMOPORIN, 2 BETA-BARREL, TRANSMEMBRANE			METAL BINDING PROTEIN BETA SANDWICH, CALCIUM-BINDING PROTEIN METAL BINDING 2
Coumpound	ASN SER); IKAP 9 CHAIN: 1; IKAP 10	ALKALINE PROTEASE; IKAP 4 CHAIN: P; IKAP 5 TETRAPEPTIDE (GLY SER ASN SER); IKAP 9 CHAIN: I; IKAP 10	ALKALINE PROTEASE; IKAP 4 CHAIN: P; IKAP 5 TETRAPEPTIDE (GLY SER ASN SER); IKAP 9 CHAIN: I; IKAP 10	ALKALINE PROTEASE; IKAP 4 CHAIN: P; IKAP 5 TETRAPEPTIDE (GLY SER ASN SER); IKAP 9 CHAIN: I; IKAP 10	OMPK36; CHAIN: A, B, C;	OMPK36; CHAIN: A, B, C;	OMPK36; CHAIN: A, B, C;	OUTER MEMBRANE PROTEIN PHOSPHOPORIN (PHOE) 1PHO 3	OUTER MEMBRANE PROTEIN PHOSPHOPORIN (PHOE) 1PHO 3	LAMININ ALPHA2 CHAIN; CHAIN: A, B, C, D;
SeqFold score										
PMF score		-0.19	-0.2	-0.19	-0.2	-0.19	-0.2	-0.2	-0.19	0.1
Verify score		1:	1.03	0.87	1.07	1.04	0.87	0.83	1.15	0.51
PSI- BLAST		5.60E-10	8.40E-14	1.10E-13	1.10E-31	1.40E-32	4.20E-27	1.40E-23	7.00E-27	2.80E-13
End AA		743	858	1244	1348	1043	1241	1045	1214	225
Start AA		482	530	890	1004	707	875	692	872	78
Chain ID		<u>a</u> .	<u>a</u>	L	¥	Ą	Α.			¥
PDB ID		lkap	lkap	1kap	losm	Iosm	losm	1pho	Ipho	1qu0
SEQ NO:		502	502	502		502	205	502	202	502

PDB annotation	PROTEIN	INTEGRAL MEMBRANE PROTEIN PORIN MATRIX PORIN, OMPF PORIN; 20MF 7 PORIN, MEMBRANE PROTEIN 20MF 12	TRANSFERASE PROTEIN-ACETYL COENZYME A COMPLEX, ACETYLTRANSFERASE	SIGNAL TRANSDUCTION PROTEIN	SIGNAL TRANSDUCTION PROTEIN	CYTOSKELETON	CYTOSKELETON	CYTOSKELETON	SIGNALING PROTEIN DAPPL PHISH	BAM32: PLECKSTRIN, 3-	PHOSPHOINOSITIDES, INOSITOL TETRAKISPHOSPHATE 2 SIGNAL	TRANSDUCTION PROTEIN, ADAPTOR PROTEIN			SIGNAL TRANSDUCTION SON OF	SEVENLESS, FLECASITAIN, SON OF SEVENLESS, SIGNAL TRANSDUCTION		ACTIN-BINDING PROTEIN ACTIN- BINDING PROTEIN, CALCIUM-
Coumpound		MATRIX PORIN OUTER MEMBRANE PROTEIN F; 20MF 5 CHAIN: NULL; 20MF 6	HPA2 HISTONE ACETYLTRANSFERASE; CHAIN: A, B, C, D;	BETA-SPECTRIN; 1BTN 4 CHAIN: NULL; 1BTN 5	BETA-SPECTRIN; 1BTN 4 CHAIN: NULL; 1BTN 5	BETA-SPECTRIN; 1DRO 6 CHAIN: NULL; 1DRO 7	BETA-SPECTRIN; IDRO 6	BETA-SPECTRIN; IDRO 6	DITAL ADAPTOR OF	PHOSPHOTYROSINE AND 3-	CHAIN: A;		PHOSPHORYLATION PLECKSTRIN (N-TERMINAL PLECKSTRIN HOMOLOGY DOMAIN) MUTANT IPLS 3 WITH LEU GLU (HIS)6	TERMINUS IPLS 4 (INS(G105- LEHHHHHH) (NMR, 25 STRUCTURES) IPLS 5	SOS 1; CHAIN: NULL;			T-FIMBRIN; CHAIN: NULL;
SeqFold score						50.76												
PMF		-0.18	0.06	98.0	0.78		0.39	0.75	-0.07	;			0.16		0.01			0.72
Verify score		1	-0.12	0.36	0.38		0.08	0.35	0.15	<u>}</u>			0.05		-0.02			-0.05
PSI- BLAST		7.00E-21	0.0028	1.70E-22	2.80E-24	7.00E-28	5.10E-17	7.00E-28	3.40F-17				1.70E-18		2.80E-14			1.20E-32
End		1202	282	181	187	192	190	189	185	}			188	_	190			250
Start		842	227	82	83	73	83	35	88	}			88		80			103
Chain			¥						\ <	:								
PDB ID		2omf	lqsm	15tn	l btn	Idro	Idro	upi	Ifao				1pls		1 pms		1	laoa
SEQ NO		502	905	207	207	207	207	207	507				507		207		[SOS

PDB annotation	BINDING, PHOSPHORYLATION	STRUCTURAL PROTEIN DYSTROPHIN, MUSCULAR DYSTROPHY, CALPONIN HOMOLOGY DOWARD, 2 ACTIN-BINDING, UTROPHIN	STRUCTURAL PROTEIN CALPONIN HOMOLOGY DOMAIN, DOMAIN SWAPPING, ACTIN BINDING, 2 UTROPHIN, DYSTROPHIN, STRUCTURAL PROTEIN	STRUCTURAL PROTEIN TWO REPEATS OF SPECTRIN, ALPHA HELICAL LINKER REGION, 2.2 TANDEM 3-HELIX COILED-COILS, STRUCTURAL PROTEIN	ENDOCYTOSIS/EXOCYTOSIS NSECI; PROTEIN-PROTEIN COMPLEX,	MULTI-SUBUNIT	ENDOCYTOSIS/EXOCYTOSIS NSECI; PROTEIN-PROTEIN COMPLEX, Mil IT-SIIRI NIT	ENDOCYTORICE NEED 1.	PROTEIN-PROTEIN COMPLEX, MULTI-SUBUNIT	ENDOCYTOSIS/EXOCYTOSIS SYNAPTOTAGMIN ASSOCIATED 35 KDA PROTEIN, P35A, THREE HELIX BUNDLE	MEMBRANE PROTEIN FOUR HELIX BUNDLE, ALPHA HELIX	CONTRACTILE PROTEIN TRIPLE. HELIX COILED COIL, CONTRACTILE PROTEIN		CHAPERONE HSP40; CHAPERONE, HEAT SHOCK, PROTEIN FOLDING, DNAK	CHAPERONE HSP40; CHAPERONE, HEAT SHOCK, PROTEIN FOLDING,
Coumpound		DYSTROPHIN; CHAIN: A, B, C, D;	UTROPHIN ACTIN BÎNDING REGION; CHAIN: A, B;	ALPHA SPECTRIN; CHAIN: A, B, C;	SYNTAXIN BINDING PROTEIN 1; CHAIN: A;	SYNTAXIN IA; CHAIN: B;	SYNTAXIN BINDING PROTEIN 1; CHAIN: A; SYNTAXIN 1A: CHAIN: R:	CYNTAXIN BINDING	PROTEIN I; CHAIN: A; SYNTAXIN 1A; CHAIN: B;	SYNTAXIN-1A; CHAIN: A, B, C,	SSO1 PROTEIN; CHAIN: A;	HUMAN SKELETAL MUSCLE ALPHA-ACTININ 2; CHAIN: A;		DNAJ; CHAIN: NULL;	DNAJ; CHAIN: NULL;
SeqFold score														57.73	
PMF		0.03	0.27	60.0	0.05		0.3	003	3	0.03	0.18	90:0			
Verify score		-0.01	0.01	-0.06	-0.13		-0.31	ڄ	}	0.25	-0.45	-0.12			0.45
PSI- BLAST		8.50E-30	1.70E-29	2.80E-09	1.40E-18		1.10E-10	1.10E-08		1.10E-09	2.80E-05	4.20E-21		6.80E-33	6.80E-33
End		248	248	562	460		261	940		933	686	467		124	122
Start AA		106	110	365	208		364	713		804	794	205		94	47
Chain ID		Ą	. ∀	V	Ø	6	ΣQ.	В.		٧	٧	4			
PDB ID		14хх	Iqag	lcun	IdnI	:	Idal	Idai		lez3	1fio	Iquu	1	1640	1bq0
SEQ NO:		208	508	515	515		cic	515		515	515	515		516	516

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PDB annotation	DNAK	MOLECULAR CHAPERONE HDJ-1; MOLECULAR CHAPERONE	MOLECULAR CHAPERONE HDI-1; MOLECULAR CHAPERONE				SUGAR BINDING PROTEIN BETA TREFOIL, MULTILECTIN RECEPTOR, PITUITARY HORMONES, 2 SULFATED CARBOHYDRATE	TRANSFERASE GLYCOSYLTRANSFERASE	TRANSFERASE GLYCOSYLTRANSFERASE	HYDROLASE XYLAN DEGRADATION				
Coumpound		HUMAN HSP40; CHAIN: NULL;	HUMAN HSP40; CHAIN: NULL;		COMPLEX (CLYCOSIDASE/CARBOHYD RATE) ABRIN-A COMPLEXED WITH TWO SUGAR CHAINS IABR 3	COMPLEX (GLYCOSIDASE/CARBOHYD RATE) ABRIN-A COMPLEXED WITH TWO SUGAR CHAINS I ABR 3	MANNOSE RECEPTOR; CHAIN: A;	SPORE COAT POLYSACCHARIDE BIOSYNTHESIS PROTEIN CHAIN: A;	SPORE COAT POLYSACCHARIDE BIOSYNTHESIS PROTEIN CHAIN: A:	ENDO-1,4-BETA-XYLANASE; CHAIN: A, B;	GLYCOSIDASE RICIN (E.C.3.2.2.2) 2AAI 3	OXYGEN TRANSPORT	HUMAN FETAL F=/11\$=) IFDHG 1 IFDHH 2	OXYGEN TRANSPORT HEMOGLOBIN (DEOXY, HUMAN FETAL F=/(1\$=) IFDHG 1 IFDHH 2
SeqFold score		52.89				,						112.26		
PMF score			-		0.01	0.04	0.13	0.11	o	0.96	0.03			1
Verify score			0.21		-0.08	0.04	-0.22	0.17	-0.1	0.08	0.07			0.29
PSI- BLAST		1.00E-30	1.00E-30		1.40E-10	6.80E-30	0.0017	5.10E-23	7.00E-45	1.70E-31	1.70E-28	1.00E-39		1.00E-39
End	L	125	122		547	547	513	333	371	548	547	92		32
Start AA	_	46	48		378	412	444	111	111	422	413	_		-
Chain					æ	EI .	V	V	٧	A	В	0		ڻ ن
PDB		1hdj	1hdj		labr	labr	Idag	lqgq	lqgq	lxyf	2aai	1fgh		1fdh
SEQ NO ID		516	516		522	522	522	522	522	522	522	523		523

PDB annotation	OXYGEN STORAGE/TRANSPORT HEMOGLOBIN, DEOXY FORM	TRANSFERASE IL-2-INDUCIBLE T- CELL KINASE; TRANSFERASE, REGULATORY INTRAMOLECULAR COMPLEX, KINASE	COMPLEX (ADAPTOR PROTEIN/PEPTIDE) ASH, GROWTH FACTOR RECEPTOR-BOUND PROTEIN 2: COMPLEX (ADAPTOR PROTEIN/PEPTIDE), SH3 DOMAIN, 2 GUANINE-NUCLEOTIDE RELEASING FACTOR	MYOSIN MYOSIN MOTOR					MYOSIN MYOSIN MOTOR				MUSCLE PROTEIN MDE; MUSCLE PROTEIN	MUSCLE PROTEIN MDE; MUSCLE PROTEIN	MUSCLE PROTEIN MUSCLE PROTEIN	MUSCLE PROTEIN MUSCLE PROTEIN	CONTRACTILE PROTEIN MYOSIN	MOTOR, CONFORMATIONAL CHANGES			
Coumpound	HEMOGLOBIN; CHAIN: A, C; HEMOGLOBIN; CHAIN: B, D;	ITK; CHAIN: NULL;	GRB2; CHAIN: A; SOS; CHAIN: B:	MYOSIN HEAVY CHAIN:	CHAIN: A; MYOSIN	REGULATORY LIGHT	CHAIN; CHAIN: Y; MYOSIN	CHAIN: Z;	MYOSIN HEAVY CHAIN;	CHAIN: A; MYOSIN	CHAIN: CHAIN: Y: MYOSIN	ESSENTIAL LIGHT CHAIN; CHAIN: Z:	MYOSIN; CHAIN: A, B, C, D, E, F, G, H;	MYOSIN; CHAIN: A, B, C, D, E, F, G, H;	MYOSIN; CHAIN: A, B, C, D, E. F.	MYOSIN; CHAIN: A, B, C, D,	MYOSIN HEAD; CHAIN: A;	MYOSIN HEAD; CHAIN: Y; MYOSIN HEAD; CHAIN: Z;	SIGNAL TRANSDUCTION	ROTEIN GROWTH FACTOR	2 (GRB2, N-TERMINAL IGBR
SeqFold score				335.35										352.11	339.22						
PMF	1	0.15	0.89						_				-			_	_		0.72		
Verify	90.0	0.45							0.34				0.59			0.44	0.01		0.45		
PSI- BLAST	7.00E-40	1.40E-18	2.80E-18		•				0				0	0	0	0	0		1.10E-19		
End	92	1088	1087	730	· }				739				708	711	629	619	739		1089		
Start	-	1018	1037	-	•		<u>. </u>		_				-	-	-	-	_		1028		
Chain	В		. ≺	4	:				¥				4	4	<	A	4		4		
PDB	lgcv	1awj	laze	147	:				1b7t		_		Ibri	lbrl	1br2	1br2	1dfk		1gbr		
SEQ BO	523	526	526	436	3				526				526	526	526	526	526		526		

			r					
PDB annotation			SIGNAL TRANSDUCTION ADAPTOR SH2, SH3 IGRI 14	·	CONTRACTILE PROTEIN MYOSIN, DICTYOSTELIUM, MOTOR, MANT, ATPASE, ACTIN-BINDING, 2 COILED COIL	CONTRACTILE PROTEIN MYOSIN, DICTYOSTELIUM, MOTOR, MANT, ATPASE, ACTIN-BINDING, 2 COILED COIL	CONTRACTILE PROTEIN ATPASE, MYOSIN, COILED COIL, ACTIN- BINDING, ATP-BINDING, 2 HEPTAD REPEAT PATTERN, METHYLATION, ALKYLATION, 3 PHOSPHORYLATION, CONTRACTILE PROTEIN	CONTRACTILE PROTEIN ATPASE, MYOSIN, COILED COIL, ACTIN- BINDING, ATP-BINDING, 2 HEPTAD REPEAT PATTERN, METHYLATION, ALKYLATION, 3 PHOSPHORYLATION, CONTRACTILE PROTEIN
Coumpound	3 SH3 DOMAIN) COMPLEXED WITH SOS-A PEPTIDE IGBR 4 (NMR, 29 STRUCTURES) IGBR 5	ADAPTOR PROTEIN CONTAINING SH2 AND SH3 GROWTH FACTOR RECEPTOR-BOUND PROTEIN 2 (GRB2) 1GFC 3 (C- TERMINAL SH3 DOMAIN) (NMR, MINIMIZED MEAN STRUCTURE) 1GFC 4	GROWTH FACTOR BOUND PROTEIN 2; 1GRI 5 CHAIN: A, B; 1GRI 6	PHOSPHORIC DIESTER HYDROLASE PHOSPHOLIPASE C-GAMMA (SH3 DOMAIN) (E.C.3.1.4.11) IHSQ 3 (NMR, MINIMIZED MEAN STRUCTURE) IHSQ 4	MYOSIN; CHAIN: NULL;	MYOSIN; CHAIN: NULL;	MYOSIN: CHAIN: NULL;	MYOSIN; CHAIN: NULL;
SeqFold score						348.12	269.23	
PMF score		0.59	0.12	1202.08	-			
Verify score		0.29	-0.23	0.24	0.18			0.37
PSI- BLAST		1.30E-19	8.40E-17	1.40E-18	0	0	o .	0
End AA		6801	1089	1089	678	619	809	809
Start		1035	896	1032	-	-	.	-
Chain ID			<					
PDB		lgfc	lgri	lhsq	Ilvk	livk	lmn d	lmn d
S B S		526	526	526	526	526	526	526

PDB annotation	CIRCULAR PERMUTANT PWT; CIRCULAR PERMUTANT, SH3 DOMAIN, CYTOSKELETON	TYROSINE-PROTEIN KINASE BRUTONS TYROSINE KINASE, B CELL PROGENITOR KINASE, TRANSFERASE, TYROSINE-PROTEIN KINASE, PHOSPHORYLATION, 2 SH3 DOMAIN	SIGNAL TRANSDUCTION PROTEIN SRC-HOMOLOGY 3 (SH3) DOMAIN, PEPTIDE-BINDING PROTEIN, ISEM 18 2 GUANINE NUCLEOTIDE EXCHANGE FACTOR ISEM 19	MUSCLE PROTEIN MUSCLE PROTEIN, MYOSIN SUBFRAGMENT-1, MYOSIN HEAD, 2 MOTOR PROTEIN	MUSCLE PROTEIN MUSCLE PROTEIN, MYOSIN SUBFRAGMENT-1, MYOSIN HEAD, 2 MOTOR PROTEIN	TRANSFERASE HCK; SH3, PROTEIN TYROSINE KINASE, SIGNAL TRANSDUCTION, 2 TRANSFERASE	ELECTRON TRANSPORT ELECTRON TRANSPORT, THIOL-DISULFIDE OXIDOREDUCTASE, 2 THIOLTRANSFERASE, THIOREDOXIN SUPERFAMILY	ELECTRON TRANSPORT ELECTRON TRANSPORT, THIOL-DISULFIDE OXIDOREDUCTASE, 2 THIOLTRANSFERASE, THIOREDOXIN SUPERFAMILY	TRANSFERASE ACETYLTRANSFERASE	TRANSFERASE AAC; AMINOGLYCOSIDE 6'-N- ACETYLTRANSFERASE, ANTIBIOTIC
Coumpound	ALPHA SPECTRIN; CHAIN: NULL;	TYROSINE-PROTEIN KINASE BTK; CHAIN: A;	SEM-5; ISEM 3 CHAIN: A, B; ISEM 5 10 RESIDUE PROLINE-RICH PEPTIDE FROM MSOS 1SEM 8 CHAIN: C, D 1SEM 10	MYOSIN; CHAIN: A, B, C;	MYOSIN; CHAIN: A, B, C;	HEMATOPOIETIC CELL KINASE; CHAIN: NULL;	GLUTAREDOXIN 3; CHAIN: NULL;	GLUTAREDOXIN 3; CHAIN: NULL;	ARYLALKYLAMINE N- ACETYLTRANSFERASE; CHAIN: A, B;	AMINOGLYCOSIDE N6- ACETYLTRANSFERASE TYPE I; CHAIN: A;
SeqFold score					267.23			-		
PMF score	-1.41	14.	0.94	_		0.18	0.68	96:0	96.0	0.95
Verify score	29.0	0.25	0.63	-0.02		0.32	-0.21	0.17	0.25	0.29
PSI- BLAST	7.00E-20	4.20E-18	4.20E-19	0	0	1.10E-18	0.0015	9.80E-06	3.40E-19	6.80E-08
End	1089	1089	1089	731	737	1089	96	88	159	184
Start AA	1032	1036	1035	-	-	1032	. 14	23	41	24
Chain D		∢	V	¥	¥				В	Y _
PDB CI	1pwt	1qfy	1sem	2mys	2mys	4hck	3grх	Здтх	1966	1881
SEQ PS PS	526	526	526	526	526		528	528	529	529

SEQ	PDB ID	Chain ID	Start	End	PSI- BLAST	Verify score	PMF score	SeqFold score	Coumpound	PDB annotation
ġ							9			2 RESISTANCE, ACETYL COENZYME A
529	1cjw	¥.	14	159	1.20E-19	0.5	0.88		SEROTONIN N- ACETYLTRANSFERASE; CHAIN: A;	TRANSFERASE N-ACETYL TRANSFERASE
529	lcm 0	Ф	36	174	2.80E-13	0.39	0.81		P300/CBP ASSOCIATING FACTOR; CHAIN: B, A;	SIGNALING PROTEIN P300/CBP ASSOCIATED FACTOR, COENZYME A, ACETYLTRANSFERASE, 2 COACTIVATOR, SIGNALING PROTEIN
529	1cm 0	m	77	184	1.70E-05	0.15	0.19		P300/CBP ASSOCIATING FACTOR; CHAIN: B, A;	SIGNALING PROTEIN P300/CBP ASSOCIATED FACTOR, COENZYME A, ACETYLTRANSFERASE, 2 COACTIVATOR, SIGNALING PROTEIN
529	lqsm	<	=	156	1.50E-13	0.13	0.33		HPA2 HISTONE ACETYLTRANSFERASE; CHAIN: A, B, C, D;	TRANSFERASE PROTEIN-ACETYL COENZYME A COMPLEX, ACETYLTRANSFERASE
529	lqsm	∢	12	165	4.20E-19	0.15	0.23		HPA2 HISTONE ACETYLTRANSFERASE; CHAIN: A, B, C, D;	TRANSFERASE PROTEIN-ACETYL COENZYME A COMPLEX, ACETYLTRANSFERASE
529	lqst	<	79	183	3.40E-07	-0.06	61.0		TGCNS HISTONE ACETYL TRANSFERASE; CHAIN: A;	TRANSFERASE HISTONE ACETYLTRANSFERASE, GCN5- RELATED N-ACETYLTRANSFERASE, 2 COA BINDING PROTEIN
529	lygh	· •		179	1.00E-05	0.09	0.04		TRANSCRIPTIONAL ACTIVATOR GCN5; CHAIN: A, B;	GENE REGULATION ADA4; TRANSCRPTIONAL REGULATION, HISTONE ACETYLATION, N- 2 ACETYLTRANSFERASE, GCN5 RELATED N-ACETYLTRANSFERASE FAMILY, 3 GENE REGULATION
534	lclg	4	9	296	1.50E-54	-0.48	-		TROPOMYOSIN; CHAIN: A, B, C, D	CONTRACTILE PROTEIN TROPOMYOSIN COILED-COIL ALPHA- HELICAL, CONTRACTILE PROTEIN
534	lclg	<	E	245	3.40E-49	-0.17	0.64		TROPOMYOSIN; CHAIN: A, B, C, D	CONTRACTILE PROTEIN TROPOMYOSIN COILED-COIL ALPHA- HELICAL, CONTRACTILE PROTEIN
534	lc1g	∢ .	m	248	5.10E-53	-0.17	0.94		TROPOMYOSIN; CHAIN: A, B, C, D	CONTRACTILE PROTEIN TROPOMYOSIN COILED-COIL ALPHA-HELICAL, CONTRACTILE PROTEIN
535	lclg	⋖	3	296	1.50E-54	-0.48			TROPOMYOSIN; CHAIN: A,	CONTRACTILE PROTEIN

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PDB annotation	TROPOMYOSIN COILED-COIL ALPHA- HELICAL, CONTRACTILE PROTEIN	CONTRACTILE PROTEIN TROPOMYOSIN COILED-COIL ALPHA-HELICAL, CONTRACTILE PROTEIN	CONTRACTILE PROTEIN TROPOMYOSIN COILED-COIL ALPHA-HELICAL, CONTRACTILE PROTEIN		COMPLEX (INHIBITOR/NUCLEASE) COMPLEX (INHIBITOR/NUCLEASE), COMPLEX (RI-ANG), HYDROLASE 2	MOLECULAR RECOGNITION,	EPITOPE MAPPING, LEUCINE-RICH 3 REPEATS	COMPLEX (NUCLEAR PROTEIN/RNA)	COMPLEX (NUCLEAR PROTEIN/RAN), RNA, SNRNP, RIBONUCLEOPROTEIN	COMPLEX (NUCLEAR PROTEIN/RNA)	COMPLEX (NUCLEAR PROTEIN/RNA),	RNA, SNRNP, RIBONUCLEOPROTEIN	COMPLEX (NUCLEAR PROTEIN/RNA)	COMPLEX (NUCLEAR PROTEIN/RNA),	RNA, SNRNP, RIBONUCLEOPROTEIN	COMPLEX (NUCLEAR PROTEIN/RNA)	COMPLEX (NUCLEAR PROTEIN/RNA),	RNA, SNRNP, RIBONUCLEOPROTEIN	COMPLEX (NUCLEAR PROTEIN/RNA)	COMPLEX (NUCLEAR PROTEIN/RNA), BNA SNBNIB BIBONI ICI EOBBOTEIN	COMPLEX ONICLEAR PROTEIN/RNA)	COMPLEX (NUCLEAR PROTEIN/RNA).	RNA, SNRNP, RIBONUCLEOPROTEIN	COMPLEX (NUCLEAR PROTEIN/RNA)	COMPLEX (NUCLEAR PROTEIN/RNA),	RNA, SNRNP, RIBONUCLEOPROTEIN	CELL ADHESION LEUCINE RICH	REPEAT, CALCIUM BINDING, CELL ADHESION	CELL ADHESION LEUCINE RICH REPEAT, CALCIUM BINDING, CELL
Coumpound	в, с, D	TROPOMYOSIN; CHAIN: A, B, C, D	TROPOMYOSIN; CHAIN: A, B, C, D		RIBONUCLEASE INHIBITOR; CHAIN: A, D; ANGIOGENIN; CHAIN: B, E;			U2 RNA HAIRPIN IV; CHAIN:	Q, R; U2 A; CHAIN: A, C; U2 B": CHAIN: B. D:	U2 RNA HAIRPIN IV; CHAIN:	Q, R; U2 A'; CHAIN: A, C; U2	B"; CHAIN: B, D;	U2 RNA HAIRPIN IV; CHAIN:	Q, R; U2 A'; CHAIN: A, C; U2	B"; CHAIN: B, D;	U2 RNA HAIRPIN IV; CHAIN:	Q, R; U2 A'; CHAIN: A, C; U2	B"; CHAIN: B, D;	U2 RNA HAIRPIN IV; CHAIN:	O, R; U2 A'; CHAIN: A, C; U2	112 RNA HAIRPIN IV. CHAIN:	O. R: U2 A': CHAIN: A. C: U2	B"; CHAIN: B, D;	U2 RNA HAIRPIN IV; CHAIN:	Q, R; U2 A'; CHAIN: A, C; U2	B"; CHAIN: B, D;	INTERNALIN B; CHAIN: A;		INTERNALIN B; CHAIN: A;
SeqFold																													
PMF score		0.64	0.94		0.47			1	•	0.45			0.29			0.12			6:0		0.29	ì		0.03			8.0		0.21
Verify score		-0.17	-0.17		0.29			0.75		0.42			0.59			-0.09			0.65		0.48	2		0.1			0.41		0.19
PSI- BLAST		3.40E-49	5.10E-53		4.20E-25			2.80E-22		4.20E-18			0.00017			3.40E-06			5.60E-23		0.00017			3.40E-06			1.70E-22		1.00E-16
End AA		245	248		306			297		306			284			<u>‡</u>			301		284	i 		144			586		350
Start AA		e	3		و			136		164			210			40			136		210	· ·		40			127		183
Chain ID		A	A		4			A		A			A			¥			ပ		U	,		၁			4	•	¥
PDB ID		lclg	lclg		1a4y			1a9n		189n			189n			1a9n			1a9n		Ia9n			1a9n			140b		1d0b
SEQ B B		535	535		538			538		538			538			538			238		538			538			538		538

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PDB annotation	ADHESION	CELL ADHESION LEUCINE RICH REPEAT, CALCIUM BINDING, CELL ADHESION	CELL ADHESION LEUCINE RICH REPEAT, CALCIUM BINDING, CELL ADHESION	CELL ADHESION LEUCINE RICH REPEAT, CALCIUM BINDING, CELL ADHESION	TRANSFERASE CRYSTAL STRUCTURE, RAB GERANYLGERANYLTRANSFERASE,	2.0 A 2 RESOLUTION, N- FORMYLMETHIONINE, ALPHA SUBUNIT, BETA SUBUNIT	CONTRACTILE PROTEIN LEUCINE- RICH REPEAT, BETA-BETA-ALPHA CYLINDER, DYNEIN, 2 CHI, AMYDOMONAS, PI, AGELLA	RNA BINDING PROTEIN TAP (NFX1); RIBONUCLEOPROTEIN (RNP,RBD OR PDAA AND I ELICATE DICE DEDEAT 2	(LRR)	RNA BINDING PROTEIN TAP (NFX1); RIBONUCLEOPROTEIN (RNP,RBD OR RRM) AND LEUCINE-RICH-REPEAT 2 (LRR)	RNA BINDING PROTEIN TAP (NFX1); RIBONUCLEOPROTEIN (RNP, RBD OR RRM) AND LEUCINE-RICH-REPEAT 2 (LRR)	LIGASE CYCLIN A/CDK2- ASSOCIATED PROTEIN P45; CYCLIN A/CDK2-ASSOCIATED PROTEIN P19; SKP1, SKP2, F-BOX, LRR, LEUCINE- RICH REPEAT, SCF, UBIQUITIN, 2 E3, THEIN INTEN DE OTTEN I ICA SE	LIGASE CYCLIN A/CDK2- ASSOCIATED PROTEIN P45; CYCLIN
Coumpound		INTERNALIN B; CHAIN: A;	INTERNALIN B; CHAIN: A;	INTERNALIN B; CHAIN: A;	RAB GERANYLGERANYLTRANSF ERASE ALPHA SUBUNIT;	CHAIN: A, C; RAB GERANYLGERANYLTRANSF ERASE BETA SUBUNIT; CHAIN: B, D;	OUTER ARM DYNEIN; CHAIN: A;	NUCLEAR RNA EXPORT FACTOR 1; CHAIN: A, B;		NUCLEAR RNA EXPORT FACTOR 1; CHAIN: A, B;	NUCLEAR RNA EXPORT FACTOR 1; CHAIN: A, B;	SKP2; CHAIN: A, C, E, G, I, K, M, O; SKP1; CHAIN: B, D, F, H, I, L, N, P;	SKP2; CHAIN: A, C, E, G, I, K, M, O; SKP1; CHAIN: B, D, F,
SeqFold score													
PMF score		0.11	0.15	0.22	-0.14		0.16	0.13		0.03	0.1	0.49	-0.09
Verify score		0.3	0.05	0.17	0.02		-0.03	-0.32		0.1	-0.41	0.63	80.0
PSI- BLAST		5.10E-24	1.00E-25	8.50E-25	1.20E-09		6.80E-13	1.00E-06		1.70E-07	1.00E-06	5.10E-13	2.80E-17
End AA		162	161	212	130		143	120		290	120	284	339
Start		E	32	57	7	-11	16	. 52		227	52	111	6
Chain ID		¥	V V	⋖	4		4	⋖		B	മ	¥	₹
PDB ID		1406	140b	140b	1dce		6sp1	1601		1601	1601	1fqv	lfqv
S B S		538	538	538	538		238	238		538	538	538	238

PDB annotation	A/CDK2-ASSOCIATED PROTEIN P19; SKP1, SKP2, F-BOX, LRR, LEUCINE- RICH REPEAT, SCF, UBIQUITIN, 2 E3, UBIQUITIN PROTEIN LIGASE	LIGASE CYCLIN A/CDK2- ASSOCIATED P45; CYCLIN A/CDK2- ASSOCIATED P19; SKP1, SKP2, F-BOX, LERS, LEUCINE-RICH REPEATS, SCF, 2 UBIQUITIN, E3, UBIQUITIN PROTEIN LIGASE	LIGASE CYCLIN A/CDK2- ASSOCIATED P45; CYCLIN A/CDK2- ASSOCIATED P19; SKP1, SKP2, F-BOX, LERS, LEUCINE-RICH REPEATS, SCF, 2 UBIQUITIN, E3, UBIQUITIN PROTEIN LIGASE	TRANSCRIPTION RNAIP; RANGAP; GTPASE-ACTIVATING PROTEIN FOR SPII, GTPASE-ACTIVATING PROTEIN, GAP, RNAIP, RANGAP, LRR, LEUCINE- 2 RICH REPEAT PROTEIN, TWINNING, HEMIHEDRAL TWINNING, MEROHEDRAL TWINNING, MEROHEDRAL	ACETYLATION RNASE INHIBITOR, RIBONUCLEASE/ANGIOGENIN INHIBITOR ACETYLATION, LEUCINE- RICH REPEATS	LIGASE TRPRS; CLASS I TRNA SYNTHETASE, AARS, INDUCED FIT, TRPRS	SIGNALING PROTEIN OXYGEN SENSOR, HISTIDINE KINASE, PAS, HIGH-RESOLUTION, TWO- 2 COMPONENT SYSTEM	TO I HOME TO ME WO I WAS A STATE OF THE PERSON OF THE PERS
	A/CDK2-4 SKP1, SKJ RICH REP UBIQUITI	LIGASE C ASSOCIA ASSOCIA LRRS, LE 2 UBIQUI LIGASE	LIGASE C ASSOCIA ASSOCIA LRRS, LE 2 UBIQUI LIGASE	TRANSCI GTPASE- SPII, GTF GAP, RN/ GAP, RN/ TWINNIN TWINNIN		LIGASE SYNTHE TRPRS	SIGNALI SENSOR, HIGH-RE COMPON	
Coumpound	H, J, L, N, P;	SKP2; CHAIN: A, C; SKP1; CHAIN: B, D;	SKP2; CHAIN: A, C; SKP1; CHAIN: B, D;	GTPASE-ACTIVATING PROTEIN RNAI_SCHPO; CHAIN: A, B;	RIBONUCLEASE INHIBITOR; CHAIN: NULL;	TRYPTOPHANYL TRNA SYNTHETASE: CHAIN: A, B, C, D, E, F;	NITROGEN FIXATION REGULATORY PROTEIN FIXL; CHAIN: A;	
SeqFold score								
PMF		0.24	0	-0.11	0.22	_	0.23	
Verify score		0.13	0.29	0.26	0.43	0.54	0.23	
PSI- BLAST		5.10E-13	1.30E-23	2.80E-22	1.40E-32	0	4.20E-17	
End.		284	306	291	311	359	371	
Start AA		111	48	98	63	35	274	
Chain		4	∢ .	⋖		<	∢	
PDB CI		162	1fs2	lyrg	2bnh	1d2r	1406	
SEQ	į.	538	538	538	538	539	540	

PDB annotation	MULTIGENE FAMILY, ACETYLATION	IRON STORAGE IRON STORAGE, DIIRON	IRON STORAGE IRON STORAGE, DIIRON	IRON STORAGE IRON STORAGE	IRON STORAGE IRON STORAGE	IRON STORAGE IRON STORAGE	ISOMERASE ISOMERASE, PPIASE	COMPLEX (ISOMERASEPEPTIDE) COMPLEX (ISOMERASEPEPTIDE), CYCLOPHILIN A, HV-1 CAPSID, 2	PSEUDO-SYMMETRY	COMPLEX (ISOMERASE/PEPTIDE) COMPLEX (ISOMERASE/PEPTIDE), CYCLOPHILIN A, HIV-1 CAPSID, 2 PSEUDO-SYMMETRY				COMPLEX (ISOMERASE/IMMUNOSUPPRESSANT) CYCLOSPORIN, ISOMERASE POTAMAGE SIGNAT ICYN 19	CI VII OI TRAIDIS STOURT ON	COMPLEX	(ISOMEKASE/IMMONOSOFFKESSANI)	ROTAMASE, SIGNAL 1CYN 19	ISOMERASE(PEPTIDYL-PROLYL CISTRANS) PEPTIDYL-PROLYL CISTRANS)
Coumpound		M FERRITIN; CHAIN: A.B.C.D.E.F.G.H.I.J.K.L.M.N.O ,P.Q.R.S.T.U,V,W,X;	M FERRITIN; CHAIN: A,B,C,D,E,F,G,H,IJ,K,L,M,N,O P,Q,R,S,T,U,V,W,X;	FERRITIN; CHAIN: NULL;	FERRITIN; CHAIN: NULL;	FERRITIN; CHAIN: NULL;	CYCLOPHILIN; CHAIN: NULL;	CYCLOPHILIN A; CHAIN: A; PEPTIDE FROM THE HIV-1 CAPSID PROTEIN; CHAIN: B;		CYCLOPHILIN A; CHAIN: A; PEPTIDE FROM THE HIV-1 CAPSID PROTEIN; CHAIN: B;	ISOMERASE(PEPTIDYL- PROLYL CIS-TRANS)	STRUCTURES) ICLH 3	ISOMERASE(PEPTIDYL- PROLYL CIS-TRANS) CYCLOPHILIN (NMR, 12 STRUCTURES) ICLH 3	CYCLOPHILIN B. ICYN 6 CHAIN: A; ICYN 7 [D- (CHOLINYL)ALA]8-	CHAIN: C; ICYN 11	CYCLOPHILIN B; ICYN 6	CHAIN: A; ICYN 7 [D-	CYCLOSPORIN; ICYN 10	CYCLOPHILIN 3; CHAIN: A;
SeqFold score		221.42		284.09			94.49	89.5			9.69			103.11					
PMF			_		_	1				1			0.46			-			-
Verify score			0.53		0.5	0.5				0.7			0.28			0.52			0.62
PSI- BLAST		5.60E-88	5.60E-88	1.40E-75	1.40E-75	1.00E-74	5.10E-29	5.10E-38		5.10E-38	8.40E-39		8.40E-39	5.10E-34		5.10E-34			1.70E-34
End		199	661	200	200	200	162	163		160	166		£91	171		160			160
Start AA		29	30	29	29	29	_	-		7	-		e e	-		2			2
Chain ID		A	٧					¥		ď				<		4			<
PDB DD		1mfr	1mfr	2fha	2fha	2fha	1a58	lawq		lawq	-1clh		1clh	Icyn		1cyn			ldy w
SEQ EQ		541	541	541	541	541	246	546		546	546		546	546		546			546

PDB annotation	TRANS ISOMERASE 3, ISOMERASE, ROTAMASE	COMPLEX (ISOMERASE/PEPTIDE) ISOMERASE, ROTAMASE, COMPLEX (ISOMERASE/PEPTIDE)	COMPLEX (ISOMERASE/PEPTIDE) ISOMERASE, ROTAMASE, COMPLEX (ISOMERASE/PEPTIDE)	PEPTIDYLPROLYL CIS-TRANS ISOMERASE; CYCLOPHILIN A, CYCLOSPORIN A, PEPTIDYL CIS- TRANS ISOMERASE	ISOMERASE USA-CYP, SNUCYP-20, CYCLOPHILIN, SNRNP, SPLICEOSOMAL	ISOMERASE USA-CYP, SNUCYP-20, CYCLOPHILIN, SNRNP, SPLICEOSOMAL			·	TRANSCRIPTION REGULATION SIGMA70; RNA POLYMERASE SIGMA FACTOR, TRANSCRIPTION REGULATION	TRANSCRIPTION REGULATION PROTO-ONCOGENE, NUCLEAR
Coumpound		CYCLOPHILIN A; CHAIN: A; SUCCINYL-ALA-PRO-ALA-P- NITROANILIDE; CHAIN: B;	CYCLOPHILIN A; CHAIN: A; SUCCINYL-ALA-PRO-ALA-P- NITROANILIDE; CHAIN: B;	CYCLOPHLIN; CHAIN: A; CYCLOSPORIN A; CHAIN: D;	SNUCYP-20; CHAIN: A;	SNUCYP-20; CHAIN: A;	COMPLEX (ISOMERASEJMMUNOSUPP RESSANT) CYCLOPHILIN C COMPLEXED WITH CYCLOSPORIN A 2RMC 3	COMPLEX (ISOMERASEJIMMUNOSUPP RESSANT) CYCLOPHILIN C COMPLEXED WITH CYCLOSPORIN A 2RMC 3	COMPLEX (ISOMERASE/IMMUNOSUPP RESSANT) CYCLOPHILIN C COMPLEXED WITH CYCLOSPORIN A 2RMC 3	RNA POLYMERASE PRIMARY SIGMA FACTOR; CHAIN: NULL;	TRANSCRIPTION FACTOR PML: CHAIN: NULL:
SeqFold score		69.67						94.18			
PMF			0.98		_	0.98	-		-	0.01	0.03
Verify score			0.47	0.56	0.57	0.31	9.0		0.39	0.33	-0.43
PSI- BLAST		7.00E-41	7.00E-41	6.80E-33	1.70E-31	2.80E-36	3.40E-31	8.40E-41	8.40臣-41	0.0028	3.40E-06
End		163	162	160	160	156	160	172	171	335	363
Start AA		2	E	7	2	7	2	2	ه	219	319
Chain		∀	V	V	A	 	Y	∢	∢		
PDB UD		1lop	1lop	Iqng	1qoi	1qoi	2ттс	2ттс	2ттс	lsig	1bor
SEQ EQ	į.	546	546	546	546	546	246	546	546	548	549

PDB Chain Start End	Start	\vdash	End		PSI-	Verify	PMF	SeqFold	Соитроипа	PDB annotation
ID AA AA BLASI	AA AA BLASI	AA BLASI	DLMSI	\dashv	31016		31012	3 332		BODIES (BODS) I EI IKEMIA 2
										BUDIES (FUDS), LEUNEMIN, 2 TRÂNSCRIPTION REGULATION
1bor 321 366 9.80E-09 -0.5	366 9.80E-09	366 9.80E-09	9.80E-09		-0.5		0.16		TRANSCRIPTION FACTOR PML; CHAIN: NULL;	TRANSCRIPTION REGULATION PROTO-ONCOGENE, NUCLEAR BODIES (PODS), LEUKEMIA, 2 TRANSCRIPTION REGULATION
1chc 321 370 3.40E-12 -0.21	370 3.40E-12	370 3.40E-12	3.40E-12		-0.21		8.0		VIRUS EQUINE HERPES VIRUS-I (C3HC4, OR RING DOMAIN) ICHC 3 (NMR, 1 STRUCTURE) ICHC 4	
1dt4 A 39 · 104 2.80E-14 0.49	39 · 104 2.80E-14	· 104 2.80E-14	2.80E-14		0.49		96.0		NEURO-ONCOLOGICAL VENTRAL ANTIGEN 1; CHAIN: A;	IMMUNE SYSTEM KH DOMAIN, ALPHA-BETA FOLD, RNA-BINDING MOTIF
1dtj A 39 104 2.80E-14 0.42	39 104 2.80E-14	104 2.80E-14	2.80E-14		0.42		0.95		RNA-BINDING NEUROONCOLOGICAL VENTRAL ANTIGEN 2; CHAIN: A, B, C, D;	IMMUNE SYSTEM KH DOMAIN, ALPHA-BETA FOLD RNA-BINDING MOTIF
1dtj B 39 106 2.80E-05 0.44	39 106 2.80E-05	106 2.80E-05	2.80E-05		0.44		0.88		RNA-BINDING NEUROONCOLOGICAL VENTRAL ANTIGEN 2; CHAIN: A, B, C, D;	IMMUNE SYSTEM KH DOMAIN, ALPHA-BETA FOLD RNA-BINDING MOTIF
1dtj C 39 104 4.20E-16 0.3	39 · 104 4.20E-16	. 104 4.20E-16	4.20E-16		0.3		0.99		RNA-BINDING NEUROONCOLOGICAL VENTRAL ANTIGEN 2; CHAIN: A, B, C, D;	IMMUNE SYSTEM KH DOMAIN, ALPHA-BETA FOLD RNA-BINDING MOTIF
1dtj D 39 104 2.80E-15 0.51	39 104 2.80E-15	104 2.80E-15	2.80E-15		0.51		0.99		RNA-BINDING NEUROONCOLOGICAL VENTRAL ANTIGEN 2; CHAIN: A, B, C, D;	IMMUNE SYSTEM KH DOMAIN, ALPHA-BETA FOLD RNA-BINDING MOTIF
lec6 A 39 109 8.40E-15 0.66	39 109 8.40E-15	109 8.40E-15	8.40E-15		0.66		-		RNA-BINDING PROTEIN NOVA-2: CHAIN: A, B; 20- MER RNA HAIRPIN; CHAIN: C, D;	RNA BINDING PROTEINRNA ASTROCYTIC NOVA-LIKE RNA- BINDING PROTEIN; KH DOMAIN, ALPHA-BETA FOLD, RNA-BINDING MOTIF, PROTEINRNA 2 STRUCTURE
1fbv A 320 370 3.40E-13 0.28	320 370 3.40E-13	370 3.40E-13	3.40E-13		0.28		0.71		SIGNAL TRANSDUCTION PROTEIN CBL; CHAIN: A; ZAP-10 PEPTIDE; CHAIN: B; UBIQUITIN-CONJUGATING ENZYME E12-18 KDA UBCH7; CHAIN: C;	LIGASE CBL, UBCH7, ZAP-70, EZ, UBIQUITIN, E3, PHOSPHORYLATION, 2 TYROSIDE KINASE, UBIQUITINATION, PROTEIN DEGRADATION,
1g25 A 317 363 0.00068 -0.37	317 363 0.00068	7 363 0.00068	0.00068	П	-0.37	1 1	0.27		CDK-ACTIVATING KINASE	METAL BINDING PROTEIN RING

PDB annotation	FINGER PROTEIN MATI; RING	METAL BINDING PROTEIN RING METAL BINDING PROTEIN RING FRICER PROTEIN MATI; RING	FINGER (USHC4) DNA-BINDING PROTEIN V(D)J DNA-BINDING PROTEIN V(D)J PROTEIN 1; RAG1, V(D)J RECOMBINATION, ANTIBODY, MAD, RNG FINGER, 2 ZINC BINUCLEAR CLUSTER, ZINC FINGER, DNA-	BINDING PROTEIN RIBONUCLEOPROTEIN RNA-BINDING PROTEIN IVIG 19		TRANSFERASE METHYLTRANSFERASE	STRUCTURAL GENOMICS HYPOTHETICAL PROTEIN,	METHANOCOCCUS JANNASCHII TRANSFERASE SAM-BINDING DOMAIN, BETA-RARREI, MIXED	ALPHA-BETA, HEXAMER, 2 DIMER	TRANSFERASE (METHYLTRANSFERASE) COMT; TRANSFERASE, METHYLTRANSFERASE, NEUROTRANSMITTER	METHYLTRANSFERASE GNMT, S- ADENOSYL-L-METHIONINE:	GLYCINE METHYLTRANSFERASE		COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC
Coumpound	ASSEMBLY FACTOR MATI; CHAIN: A:	CDK-ACTIVATING KINASE ASSEMBLY FACTOR MATI, CHAIN: A.	RAGI; CHAIN: NULL;	VIGILIN; 1VIG 5 CHAIN: NULL; 1VIG 6		GLYCINE N- METHYLTRANSFERASE; CHAIN: A B C D.		\SE;		SE;	GLYCINE N- METHYLTRANSFERASE;		Cosp 2 Price Brices		QGSR ZINC FINGER QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE
SeqFold score															
PMF score		0.25	0.19	-		0.29	_	0.53			90.0		0.01		10.0
Verify score		-0.18	0.16	0.69		-0.04	0.38	-0.12	7	-0.07	-0.04		-0.28		-0.43
PSI- BLAST		1.40E-09	5.10E-12	4.20E-16		3.40E-15	8.50E-09	5.60E-07	4 200	4.20E-09	5.10E-16		1.70E-23		8.50E-25
End AA		365	363	104		163	166	164	100	100	165		490		547
Start AA		321	312	36		2	51	34	87	9	28		410		446
Chain ID		∢ .					∀	1			<		· ∀		
PDB ID		1g25	1md	lvig	157		Idus	1869	1 vid		Ixva		lalh /		lalh A
S B S		549	549	549	Ş	3	233	553	553		g	7	554		554

End PSI- BLAST Verify score PMF score SeqFold 1032 1.00E-10 0.14 -0.19 470 1.70E-12 0.27 0.52
\$.10E-09 -0.21
470 3.40E-13 -0.03 0.04
472 5.10E-13 0.04 0.24
553 1.70E-15 -0.35 0
472 2.80E-11 0.01 0.89
295 3.40E-15 -0.15 0
352 1.70E-11 0.11 -0.13

								· -	т —	_	_	_		-	
PDB annotation	TRANSDUCIN BETA SUBUNIT; GAMMAI, TRANSDUCIN GAMMA SUBUNIT; COMPLEX (GTP- BINDING/TRANSDUCER), G PROTEIN, HETEROTRIMER 2 SIGNAL TRANSDUCTION	REPLICATION DNA NUCLEOTIDE	EXCISION KEPAIK, UVKABC, HELICASE, 2 HYPERTHERMOSTABLE PROTEIN	HYDROLASE UVRB; MULTIDOMAIN PROTEIN	GENE REGULATION APO PROTEIN	GENE REGULATION APO PROTEIN	TRANSLATION YEAST INITIATION FACTOR 44, EIF44; HELICASE, INITIATION FACTOR 44, DEAD-BOX PROTEIN	TRANSLATION EUKARYOTIC INITIATION FACTOR 44; 154A, HELICASE, DEAD-BOX PROTEIN	TRANSLATION EUKARYOTIC INITIATION FACTOR 4A; 1F4A, HELICASE, DEAD-BOX PROTEIN	HELICASE HELICASE, RNA, HEPATITIS. HCV. ATPASE, NTPASE	HELICASE HELICASE, RNA, HEPATITIS, HCV. ATPASE, NTPASE	HELICASE HELICASE, RNA, HEPATITIS HCV. ATPASE, NTPASE	GENE REGULATION EIF4A; TRANSLATION INITIATION, SACCHAROMYCES CEREVISIAE, DEAD BOY 2 DECTENT FAART V	DEAD BOA LINGIBIN FAMILI	TRANSFERASE
Coumpound	BETA; CHAIN: B; GT- GAMMA; CHAIN: G;	DNA NUCLEOTIDE	EXCISION KEPAIR ENZYME UVRB; CHAIN: A;	EXCINUCLEASE ABC SUBUNIT B; CHAIN: A;	EXCINUCLEASE UVRABC COMPONENT UVRB; CHAIN: A;	EXCINUCLEASE UVRABC COMPONENT UVRB; CHAIN: A:	EUKARYOTIC INITIATION FACTOR 4A; CHAIN: A;	YEAST INITIATION FACTOR 4A; CHAIN: A, B;	YEAST INITIATION FACTOR 4A; CHAIN: A, B;	HCV HELICASE; CHAIN: A, B;	HCV HELICASE; CHAIN: A, B;	HCV HELICASE; CHAIN: A,	TRANSLATION INITIATION FACTOR 4A; CHAIN: A;		GLYCINE N-
SeqFold score									,						
PMF		0.59		0.95	0.64	0.87			_	0.19	0	0.05			0.95
Verify score		0.1		0.01	0.06	-0.03	0.8	0.74	0.64	0.39	-0.35	-0.11	0.84		0.27
PSI- BLAST		3.40E-14		3.40E-14	8.40E.47	1.50E-16	8.50E-45	1.70E-55	0	1.70E-06	5.60E-09	1.70E-06	1.70E-52		1.40E-18
End AA		317		317	379	317	374	204	374	310	316	310	203		195
Start		175		175	123	175	213	1	-	236	40	236	1		65
Chain D		A		A	A	Ą	A	¥	В	A	A	В	¥		¥
PDB ID		1640		1d2 m	×6p1	149x	1fûk	1 fuu	1 fuu	Ihei	Ihei	lhei ·	1qde		1d2h
SEQ ID NO:		557		557	557	557	557	557	557	557	557	557	257		558

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PDB annotation	METHYLTRANSFERASE	TRANSFERASE METHYLTRANSFERASE	STRUCTURAL GENOMICS HYPOTHETICAL PROTEIN, METHANOCOCCUS JANNASCHII	STRUCTURAL GENOMICS HYPOTHETICAL PROTEIN, METHANOCOCCUS JANNASCHII	TRANSFERASE SAM-BINDING DOMAIN, BETA-BARREL, MIXED ALPHA-BETA, HEXAMER, 2 DIMER	METHYLTRANSFERASE GNMT, S- ADENOSYL-L-METHIONINE: GLYCINE METHYLTRANSFERASE	METHYLTRANSFERASE ERMAM; METHYLTRANSFERASE, ERM, ERMAM, MLS ANTIBIOTICS, NMR, 2 RRNA	METHYLTRANSPERASE TRANSFERASE, METHYLTRANSFERASE, RESTRICTION SYSTEM	HYDROLASE PROTEIN-TYROSINE PHOSPHATASE; HYDROLASE, CATOLINI TYROSINE PHOSPHATASE, CATALYTIC DOMAIN, 2 WPD LOOP, SH2 DOMAIN	HYDROLASE DUAL SPECIFICITY PHOSPHATASE, MAP KINASE HYDROLASE	HYDROLASE DUAL SPECIFICITY PHOSPHATASE, MAP KINASE HYDROLASE	HYDROLASE VHR; HYDROLASE, PROTEIN DUAL-SPECIFICITY PHOSPHATASE	HYDROLASE VHR; HYDROLASE,
Coumpound	METHYLTRANSFERASE; CHAIN: A, B, C, D;	GLYCINE N- METHYLTRANSFERASE; CHAIN: A, B, C, D;	MJ0882; CHAIN: A;	MJ0882; CHAIN: A;	HNRNP ARGININE N- METHYL TRANSFERASE; CHAIN: 1, 2, 3, 4, 5, 6;	GLYCINE N- METHYLTRANSFERASE; CHAIN: A, B;	RRNA METHYLTRANSFERASE; CHAIN: NULL;	ADENINE-NG-DNA- METHYLTRANSFERASE TAQI; CHAIN: A, B;	SHP-1; CHAIN: NULL;	PYST1; CHAIN: NULL;	PYST1: CHAIN: NULL;	HUMAN VHI-RELATED DUAL-SPECIFICITY PHOSPHATASE CHAIN: A. B:	HUMAN VHI-RELATED
SeqFold score									•	137.8		96.19	
PMF		0.19	0.89	0.93	0.31	0.72	90.0	0.48	0.25		_		
Verify		-0.17	0.35	0.25	0.11	-0.08	0.44	0.09	-0.17		0.88		0.78
PSI- BLAST		2.80E-13	1.40E-12	2.80E-14	1.70E-26	· 5.10E-21	1.20E-06	1.20E-08	4.20E-05	3.40E-39	3.40E-39	1.10E-34	1.10E-34
End		223	187	213		195	157	189	301	298	287	307	298
Start		29	89	83	43	43	71	62	225	158	. 651	135	142
Chain TD		4	4	٧	-	V	,	A				¥	A
PDB UD		1d2h	1dus	1dus	1g6q	lxva	lyub	2ad m	lgwz	lmk P	Imk p	lvhr	1vhr
SEQ NO.		858	558	828	558	858	558	558	559	529	559.	529	559

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PDB annotation	PROTEIN DUAL-SPECIFICITY PHOSPHATASE	HYDROLASE VHR; HYDROLASE, PROTEIN DUAL-SPECIFICITY PHOSPHATASE		COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC	FINGER, DINA-BUNDING FROIEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC	FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGERDINA), ZINC FINGER, DNA-BINDING PROTEIN		COMPLEX (ZINC FINGER/DNA) ZINC	FINGER, PROTEIN-DNA	CRYSTAL STRUCTURE, COMPLEX	COMPLEX (ZINC FINGER/DNA) ZINC	FINGER, PROTEIN-DNA	INTERACTION, PROTEIN DESIGN, 2	CRYSTAL STRUCTURE, COMPLEX	COMPLEX (ZINC FINGER/DNA) ZINC	FINGER, PROTEIN-DNA	INTERACTION, PROTEIN DESIGN, 2	CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC	FINGER, PROTEIN-DNA	INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX
Coumpound	DUAL-SPECIFICITY PHOSPHATASE CHAIN: A, B;	HUMAN VHI-RELATED DUAL-SPECIFICITY PHOSPHATASE CHAIN: A, B;		QGSR ZINC FINGER PEPTIDE, CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX	BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE: CHAIN: A: DUPLEX	OLIGONUCLEOTIDE BINDING SITE: CHAIN: B. C.	QGSR ZINC FINGER	PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE	BINDING SITE; CHAIN: B, C,	DNA; CHAIN: A, B, D, E;	CONSENSUS ZINC FINGER	FROIEIN; CHAIN: C, F, G;	DNA-CHAIN A B D E	CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;		DNA; CHAIN: A, B, D, E;	CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;		DNA; CHAIN: A, B, D, E;	CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;
SegFold							56.74												69.14						
PMF		1		0.77	0.83				98.0			69.0			_	•							0.46		
Verify score		0.29		0.4	0.03				-0.45			0.07			80 0	3							-0.37		
PSI- BLAST		1.00E-32		6 _. 80E-22	1.50E-28	t	1.50E-28		1.20E-24			6.80E-36			\$ 10E 40	7.704.5			5.10E-49				3.40E-40		
End		282		248	276		278		288			246			276	2			277				288		
Start		146		188	194		194		222			187			103	3			193				221		
Chain		A		V	4		¥		4			ပ			ر)			U				ပ		
PDB		lvhr		lalh	lalh		lath		lalh			Ішс	>		130) ×			Ime	>			l E	>	
SEQ B SEQ		559		566	995		999		566			995			4,46	3			999				999		

PDB annotation	(ZINC FINGER/DNA)			COMPLEX (TRANSCRIPTION REGULATION/DNA) TFIIIA: 5S GENE: NMR, TFIIIA, PROTEIN, DNA, TRANSCRIPTION FACTOR, 5S RNA 2 GENE, DNA BINDING PROTEIN, ZINC FINGER, COMPLEX 3 (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION E; REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	A,	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX CTRANSCRIPTION REGULATION/DNA)	``
Coumpound		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	TRANSCRIPTION FACTOR IIIA; CHAIN: A; 58 RNA GENE; CHAIN: E, F;	TRANSCRIPTION FACTOR IIIA; CHAIN: A; 5S RNA GENE; CHAIN: E, F;	TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	YY I; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	
SeqFold score			50.62	-		56.56		
PMF score		0.92		0.01	0.16		0.89	
Verify		-0.03		-0.15	-0.28		-0.32	
PSI- BLAST		1.70B-13	6.80E-18	6.80E-18	1.40E-20	6.80E-29	6.80E-29	
End		246	280	276	277	277	276	
Start		219	193	194	198	167	186	
Chain		5	4	∢	∢	υ	U	
PDB		1me y	143	1143	1116	lubd	1 ubd	
SEQ S B S		995	996	566	566	566	999	

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PDB annotation	REGULATION/DNA) YING-YANG I; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	SCAFFOLD PROTEIN SCAFFOLD PROTEIN, PP24, PHOSPHORYLATION, HEAT DEDEAT	SCAFFOLD PROTEIN SCAFFOLD PROTEIN, PP2A, PHOSPHORYLATION, HEAT REPEAT	TRANSPORT PROTEIN SERINE-RICH RNA POLYMERASE I SUPPRESSOR PROTEIN; ARM REPEAT	TRANSPORT PROTEIN SERINE-RICH RNA POLYMERASE I SUPPRESSOR PROTEIN; ARM REPEAT	TRANSPORT PROTEIN SERINE-RICH RNA POLYMERASE I SUPPRESSOR PROTEIN; ARM REPEAT	TRANSPORT PROTEIN SERINE-RICH RNA POLYMERASE I SUPPRESSOR PROTEIN; ARM REPEAT	NUCLEAR IMPORT RECEPTOR KARYOPHERIN ALPHA; NUCLEAR IMPORT RECEPTOR, NUCLEAR LOCALIZATION SIGNAL, 2
Coumpound	ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	PROTEIN PHOSPHATASE PP2A; CHAIN: A, B;	PROTEIN PHOSPHATASE PP2A; CHAIN: A, B;	KARYOPHERIN ALPHA; CHAIN: A, B; MYC PROTO- ONCOGENE PROTEIN; CHAIN: C, D, E, F.	KARYOPHERIN ALPHA; CHAIN: A. B. MYC PROTO- ONCOGENE PROTEIN; CHAIN: C. D. E. F:	KARYOPHERIN ALPHA; CHAIN: A, B; MYC PROTO- ONCOGENE PROTEIN; CHAIN: C. D. P. P.	KARYOPHERIN ALPHA; CHAIN: A, B, MYC PROTO- ONCOGENE PROTEIN; CHAIN: C, D, E, F:	IMPORTIN ALPHA; CHAIN: . A;
SeqFold score		57.42		161.25						
PMF			0.72		0.71	-	0.83	_	-	0.96
Verify score			0.23		-0.03	0.42	-0.06	0.44	0.34	0.38
PSI- BLAST		1.50E-26	1.50E-26	1.70E-40	1.70E-40	1.30E-43	1.40E-31	1.70E-37	3.40E-28	5.10E-21
End		278	278	909	605	602	459	109	390	603
Start		133	188	1	6	144	81	184	7	304
Chain		∢	A	A A	٧	¥	A	∢	∢ .	∢
PDB UD		2gli	2gli	1b3u	163u	1ee4	lee4	1ee4	1cc4	lial
SEQ US	2	999	266	172	571	122	571	571	571	571

PDB annotation	ARMADILLO REPEATS, AUTOINHIBITION, INTRASTERIC REGULATION	NUCLEAR IMPORT RECEPTOR KARYOPHERIN ALPHA; NUCLEAR IMPORT RECEPTOR, NUCLEAR CACALIZATION SIGNAL, 2 ARMADILLO REPEATS, AUTONHIBITION, INTRASTERIC REGULATION	NUCLEAR IMPORT RECEPTOR KARYOPHERIN ALPHA; NUCLEAR IMPORT RECEPTOR, NUCLEAR LOCALIZATION SIGNAL, 2 ARMADILLO REPEATS, AUTONHIBITION, INTRASTERIC REGULATION	NUCLEAR IMPORT RECEPTOR KARYOPHERIN ALPHA; NUCLEAR IMPORT RECEPTOR, NUCLEAR LOCALIZATION SIGNAL, 2 ARMADILLO REPEATS, AUTONHIBITION, INTRASTERIC REGULATION	SMALL GTPASE KARYOPHERIN BETA, P95 SMALL GTPASE, NUCLEAR TRANSPORT RECEPTOR	SMALL GTPASE KARYOPHERIN BETA, P95 SMALL GTPASE. NUCLEAR TRANSPORT RECEPTOR	NUCLEAR TRANSPORT PROTEIN COMPLEX HEAT REPEATS, NUCLEAR TRANSPORT PROTEIN COMPLEX	TRANSPORT RECEPTOR KARYOPHERIN BETA-1, NUCLEAR FACTOR P97, IMPORTIN IMPORTIN ALPHA-2 SUBUNIT, KARYOPHERIN ALPHA-2 TRANSPORT RECEPTOR, NUCLEAR IMPORT, HEAT MOTIF, NLS-BINDING	STRUCTURAL PROTEIN ARMADILLO
	ARMA AUTO REGU	NUCL KARY IMPOI LOCA ARM/ AUTO	NUCL KARY IMPO LOCA ARMA AUTO REGU	NUCI KARS IMPO LOCA ARM AUTO	SMAI BETA TRAN	SMAJ BETA TRAN	COM	KAR KAR FACT ALPH ALPH NUCI NUCI	STRL
Coumpound		IMPORTIN ALPHA; CHAIN: A;	IMPORTIN ALPHA; CHAIN: A;	IMPORTIN ALPHA; CHAIN: A;	RAN; CHAIN: A, C; IMPORTIN BETA SUBUNIT; CHAIN: B, D;	RAN; CHAIN: A, C; IMPORTIN BETA SUBUNIT; CHAIN: B, D;	KARYOPHERIN BETA2; CHAIN: B; RAN; CHAIN: C;	IMPORTIN BETA SUBUNIT; CHAIN: A; IMPORTIN ALPHA-2 SUBUNIT; CHAIN: B;	BETA-CATENIN; CHAIN:
SeqFold score		152.5							
PMF			0.19	88.	9.0	-0.09	0.74	0.16	9.64
Verify			0.26	0.18	-0.01	0.11	10.0	-0.16	0.29
PSI- BLAST		1.40E-32	1.70E-09	1.40E-32	6.80E-13	5.10E-12	3.40E-49	6.80E-23	1.40E-22
End		475	607	390	909	26	605	474	200
Start		33	530	r	225	Е	9	m	8
Chain		<	<	∢	m	В	Ø	∢	
PDB		lial	lial	lial	libr	1ibr	1qbk	lqgr	2bct
SEQ	Ž	571	571	571	175	571	571	571	571

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=		KMADII	RMADII	KMADII	ARMADILLO REPEAT AKMADILLO REPEAT, BETA-CATENIN, CYTOSKELETON	ARMADILLO REPEAT ARMADILLO REPEAT, BETA-CATENIN, CYTOSKELETON	ARMADILLO REPEAT ARMADILLO REPEAT, BETA-CATENIN, CYTOSKELETON	ARMADILLO REPEAT ARMADILLO REPEAT, BETA-CATENIN, CYTOSKELETON		LIPID TRANSPORT APO A-I; LIPOPROTEIN, LIPID TRANSPORT, CHOLESTEROL METABOLISM, 2 ATHEROSCLEROSIS, HDL, LCAT- ACTIVATION	STRUCTURAL PROTEIN TWO REPEATS OF SPECTRIN, ALPHA HELICAL LINKER REGION, 2 2 TANDEM 3-HELIX COILED-COILS, STRUCTURAL PROTEIN	CONTRACTILE PROTEIN TRIPLE- HELIX COILED COIL, CONTRACTILE PROTEIN	TRANSCRIPTION REGULATION SIGMA70; RNA POLYMERASE SIGMA FACTOR, TRANSCRIPTION REGULATION
PDB annotation	ATENIN	ATENIA	ATENIN OTEIN	ATENIN	EAT AR	PEAT AR	PEAT AR ATENIN	ARMADILLO REPEAT ARI REPEAT, BETA-CATENIN, CYTOSKELETON		LIPID TRANSPORT APO A-I; CHOLESTEROL METABOLISM, 2 ATHEROSCLEROSIS, HDL, LCATACTIVATION	STRUCTURAL PROTEIN TWO REPEATS OF SPECTRIN, ALPHA HELICAL LINKER REGION, 2 2 TANDEM 3-HELIX COILED-COII STRUCTURAL PROTEIN	ROTEIN COIL, CO	TRANSCRIPTION REGULATION SIGMA70; RNA POLYMERASE S FACTOR, TRANSCRIPTION REGULATION
PDB	BETA-C	BETA-C JRAL PR	JRAL PR BETA-C JRAL PR	DRAL PR BETA-C URAL PF	LLO REI BETA-C ELETON	LLO RE BETA-C ELETON	ILLO RE BETA-C ELETON	ILLO RE BETA-C ELETON		CANSPOI OTEIN, L TEROL I SCLERC TION	STRUCTURAL PROTEIN REPEATS OF SPECTRIN, HELICAL LINKER REGIG TANDEM 3-HELIX COIL.I STRUCTURAL PROTEIN	ACTILE F	RIPTION OF RNA P
	REPEAT, BETA-CATENIN, STRUCTURAL PROTEIN	STRUCTURAL PROTEIN ARMADILLO REPEAT, BETA-CATENIN, STRUCTURAL PROTEIN	STRUCTURAL PROTEIN ARMADILLO REPEAT, BETA-CATENIN, STRUCTURAL PROTEIN	STRUCTURAL PROTEIN ARMADILLU REPEAT, BETA-CATENIN, STRUCTURAL PROTEIN	ARMADILLO REPEAT AR REPEAT, BETA-CATENIN CYTOSKELETON	ARMADILLO REPEAT ARY REPEAT, BETA-CATENIN, CYTOSKELETON	ARMADILLO REPEAT ARI REPEAT, BETA-CATENIN, CYTOSKELETON	ARMADILLO REP REPEAT, BETA-CA CYTOSKELETON		LIPID TRANSI LIPOPROTEIN CHOLESTERO ATHEROSCLE ACTIVATION	STRUCTURAL PROTEIN TWO REPEATS OF SPECTRIN, ALP HELICAL LINKER REGION, 2: TANDEM 3-HELIX COILED-CO STRUCTURAL PROTEIN	CONTRAC HELIX CC PROTEIN	TRANSCRIPTIC SIGMA70; RN/FFACTOR, TRAIREGULATION
		÷ ·	ÿ	"	"	ż	ż	ż			AIN:	USCLE IAIN:	TOR;
Coumpound		BETA-CATENIN; CHAIN: NULL;	BETA-CATENIN; CHAIN: NULL;	BETA-CATENIN; CHAIN: NULL;	BETA-CATENIN; CHAIN: NULL;	BETA-CATENIN; CHAIN: NULL;	BETA-CATENIN; CHAIN: NULL;	BETA-CATENIN; CHAIN: NULL;		APOLIPOPROTEIN A-1; CHAIN: A, B, C, D;	ALPHA SPECTRIN; CHAIN: A, B, C;	HUMAN SKELETAL MUSCLE ALPHA-ACTININ 2; CHAIN: A;	RNA POLYMERASE PRIMARY SIGMA FACTOR; CHAIN: NULL;
Coum		CATENI	CATENI	CATENI	CATEN	CATEN	-CATEN	CATEN		APOLIPOPROTEIN CHAIN: A, B, C, D;	A SPECT	AN SKEI A-ACTIN	RNA POLYMERASE PRIMARY SIGMA F/ CHAIN: NULL;
	NULL;	BETA-	BETA- 'NULL;	BETA- NULL;	BETA-	BETA-(NULĻ;	BETA-(NULL;	BETA- NULL;		CHAI	ALPHA A, B, C,	HUM/ ALPH A:	RNA J PRIM CHAIJ
SeqFold		160.13				141.33				67.34	62.14	67.92	
PMF			-	0.94	-1.41		1202.08	-					0.07
Verify score			0.39	0.31	0.57		0.45	0.35			•		-0.28
PSI- BLAST		6.80E-44	8.40E-25	6.80E-44	6.80E-18	1.40E-31	5.10E-36	1.40E-31		5.60E-11	2.80E-12	1.40E-13	2.80E-09
End AA B					386 6.8	476 1.4	604 5.1	430 1.4	_	219 5.0	219 2.1	245 1.	191 2.3
		607	431	607	3	4		4.		2	2	7	-
Star	ļ	88	6	16		13	138	51	_	ES .	9	8	31
Chain										4	∢	∢	
PDB CD		2bct	2bct	2bct	3bct	3bct	3bct	3bct		lavl	lcun	1quu	lsig
SEQ D		571	571	57.1	571	571	571	571		572	572	572	572

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PDB annotation		LIPID TRANSPORT APO A-1; LIPOPROTEIN, LIPID TRANSPORT, ATHEROSCLEROSIS, HDL, LCAT- ACTIVATION	TRANSMEMBRANE PROTEIN COLICIN, BACTERIOCIN, ION CHANNEL FORMATION, TRANSMEMBRANE 2 PROTEIN	STRUCTURAL PROTEIN TWO REPEATS OF SPECTRIN, ALPHA HELICAL LINKER REGION, 2.2 TANDEM 3-HELIX COILED-COILS, STRUCTURAL PROTEIN	STRUCTURAL PROTEIN TWO REPEATS OF SPECTRIN, ALPHA HELICAL LINKER REGION, 2.2 TANDEM 3-HELIX COILED-COILS, STRUCTURAL PROTEIN	SIGNALING PROTEIN GUANINE NUCLEOTIDE- BINDING PROTEIN 1; NUTERPERON INDUCED, DYNAMIN, RELATED, LARGE GTPASE FAMILY, SIGNALING PROTEIN	ENDOCYTOSIS/EXOCYTOSIS NSEC!; PROTEIN-PROTEIN COMPLEX. MULTI-SUBUNIT	ENDOCYTOSIS/EXOCYTOSIS NSECI; PROTEIN-PROTEIN COMPLEX, MULTI-SUBUNIT	ENDOCYTOSIS/EXOCYTOSIS SYNAPTOTAGMIN ASSOCIATED 35 KDA PROTEIN, P35A, THREE HELIX BUNDLE	SIGNALING PROTEIN GBP, GTP HYDROLYSIS, GDP, GMP, INTERFERON INDUCED, DYNAMIN 2 RELATED, LARGE GTPASE FAMILY. GMPPNP, GPPNHP.
Coumpound		APOLIPOPROTEIN A-1; CHAIN: A, B, C, D;	COLICIN 1A; CHAIN: NULL;	ALPHA SPECTRIN; CHAIN: A, B, C;	A, B, C;	INTERFERON-INDUCED GUANYLATE-BINDING PROTEIN I; CHAIN: A;	SYNTAXIN BINDING PROTEIN 1; CHAIN: A; SYNTAXIN 1A; CHAIN: B;	SYNTAXIN BINDING PROTEIN 1; CHAIN: A; SYNTAXIN 1A; CHAIN: B;	SYNTAXIN-1A; CHAIN: A, B, C,	INTERFERON-INDUCED GUANYLATE-BINDING PROTEIN 1; CHAIN: A;
SeqFold score				·			-			
PMF		-0.18	-0.2	-0.01	-0.12		-0.17	-0.19	-0.18	-0.19
Verify score		0.52	90:0	0.5	0.57	0.43	0.39	0.31	0.43	0.29
PSI- BLAST		7.00E-13	4.20E-16	5.60E-12	7.00E-15	1.40E-10	8.40E-14	1.40E-11	1.10E-16	4.20E-11
End		137	175	175	125	140	145	175	129	145
Start			7		9	16	4	&	14	11
Chain		∢		4	∢	4	æ	æ	∢	<
PDB TD	T	lavi	Icii	Icun	Icun	14g3	ldnl	1dn1	1ez3	15n
SEQ EQ	2	573	573	573	573	573	573	573	573	573

PDB annotation	PROTEIN TRANSPORT HELIX-TURN- HELIX TPR-LIKE REPEAT, PROTEIN TRANSPORT	CONTRACTILE PROTEIN TRIPLE- HELIX COILED COIL, CONTRACTILE PROTEIN	CONTRACTILE PROTEIN TRIPLE- HELIX COILED COIL, CONTRACTILE PROTEIN	CONTRACTILE PROTEIN TRIPLE- HELIX COILED COIL, CONTRACTILE PROTEIN	COMPLEX (TRANSDUCERTRANSDUCTION) GT BETA-GAMMA; MEKA, PP3; PHOSDUCIN, TRANSDUCTION, BETA- GAMMA, SIGNAL TRANSDUCTION, 2 REGULATION, PHOSPHORYLATION, G PROTEINS, THOREDOXIN, 3 VISION, MEKA, COMPLEX (TRANSDUCER/TRANSDUCTION)		TOXIN BINDING PROTEIN TWO DOMAINS: BETA PROPELLER AND ALPHA/BETA FOLD	TOXIN BINDING PROTEIN TWO DOMAINS: BETA PROPELLER AND ALPHA/BETA FOLD	TOXIN BINDING PROTEIN TWO DOMAINS: BETA PROPELLER AND ALPHA/BETA FOLD	TOXIN BINDING PROTEIN TWO DOMAINS: BETA PROPELLER AND ALPHA/BETA FOLD	TOXIN BINDING PROTEIN TWO DOMAINS: BETA PROPELLER AND ALPHA/BETA FOLD	TRANSCRIPTION INHIBITOR BETA- PROPELLER	TRANSCRIPTION INHIBITOR BETA-
Coumpound	VESICULAR TRANSPORT PROTEIN SEC17; CHAIN: A;	HUMAN SKELETAL MUSCLE ALPHA-ACTININ 2; CHAIN: A;	HUMAN SKELETAL MUSCLE ALPHA-ACTININ 2; CHAIN: A;	HUMAN SKELETAL MUSCLE ALPHA-ACTININ 2; CHAIN: A;	TRANSDUCIN; CHAIN: B. G; PHOSDUCIN; CHAIN: P;		TOLB PROTEIN, CHAIN: A,	TOLB PROTEIN; CHAIN: A;	TOLB PROTEIN; CHAIN: A;	TOLB PROTEIN; CHAIN: A;	TOLB PROTEIN; CHAIN: A;	TRANSCRIPTIONAL REPRESSOR TUP I; CHAIN: A, B. C:	TRANSCRIPTIONAL
SeqFold score			53.17										
PMF score	-0.2	-0.2		-0.18	-0.19		0.22	0.1	0	0.22	-0.12	-0.14	I
Verify score	0.2	0.4		0.36	0.08		0.47	0.37	0.41	-0.14	0.22	0.14	8.0
PSI- BLAST	1.40E-12	5.60E-12	1.40E-19	1.40E-19	8.40E-11		9.80E-18	5.60E-16	.4.20E-15	5.10E-05	1.40E-09	5.60E-91	1.70E-76
End	146	101	250	174	144		358	466	484	476	209	484	484
Start	9		9 .	٦		٠	011		797	353	. 16	601	193
Chain	V	V	∢	4	a.		A	Ą	Ą	٧	V	₹	A
PDB	lqqe	1quu	lquu	1quu	2trc		lcrz	lcrz	lcrz	lcrz	lcrz	lerj	lerj
SEQ Signal	573	573	573	573	573		574	574	574	574	574	574	574

PDB annotation	PROPELLER	TRANSCRIPTION INHIBITOR BETA- PROPELLER	COMPLEX (GTP-BINDINGTRANSDUCER) BETA1, TRANSDUCIN BETA SUBUNIT; GAMMA1, TRANSDUCIN GAMMA SUBUNIT; COMPLEX (GTP-BINDINGTRANSDUCER), G PROTEIN, HETEROTRIMER 2 SIGNAL TRANSDUCTION	COMPLEX (GTP- BINDINGTRANSDUCER) BETA1, TRANSDUCIN BETA SUBUNIT; GAMMA1, TRANSDUCIN GAMMA SUBUNIT; COMPLEX (GTP- BINDINGTRANSDUCER), G PROTEIN, HETEROTRIMER 2 SIGNAL TRANSDUCTION	COMPLEX (GTP- BNDING/TRANSDUCER) BETA1, TRANSDUCIN BETA SUBUNIT; GAMMA1, TRANSDUCIN GAMMA SUBUNIT; COMPLEX (GTP- BINDING/TRANSDUCER), G PROTEIN, HETEROTRIMER 2 SIGNAL TRANSDUCTION	COMPLEX (GTP-BINDINGTRANSDUCER) BETA1, TRANSDUCIN BETA SUBUNIT; TRANSDUCIN BETA1, TRANSDUCIN GAMMA SUBUNIT; COMPLEX (GTP-BINDINGTRANSDUCER), G PROTEIN, HETEROTRIMER 2 SIGNAL TRANSDUCTION	COMPLEX (GTP- BINDING/TRANSDUCER) BETA1, TRANSDUCIN BETA SUBUNIT; GAMMA1, TRANSDUCIN GAMMA
Coumpound	REPRESSOR TUP1; CHAIN: A, B, C;	TRANSCRIPTIONAL REPRESSOR TUP1; CHAIN: A, B, C;	GT-ALPHA/GI-ALPHA CHIMERA; CHAIN: A; GT- BETA; CHAIN: B; GT- GAMMA; CHAIN: G;	GT-ALPHA/GI-ALPHA CHIMERA; CHAIN: A; GT- BETA; CHAIN: B; GT- GAMMA; CHAIN: G;	GT-ALPHA/GI-ALPHA CHIMERA; CHAIN: A; GT- BETA; CHAIN: B; GT- GAMMA; CHAIN: G;	GT-ALPHA/GI-ALPHA CHIMERA; CHAIN: A; GT- BETA; CHAIN: B; GT- GAMMA; CHAIN: G;	GT-ALPHA/GI-ALPHA CHIMERA; CHAIN: A; GT- BETA; CHAIN: B; GT- GAMMA; CHAIN: G;
SeqFold score					•		150.13
PMF		_	1	-	-	66:0	
Verify score		0.46	0.95	0.7	0.79	0.43	·
PSI- BLAST		1.70E-67	3.40E-71	1.00E-50	6.80E- <i>79</i>	3.40E-55	3.40E-71
End		356	400	273	483	315	400
Start AA		48	101	13	190	45	59
Chain		V	В	B	æ	м	Ø
PDB D		lerj	lgot	1got	1got	lgot	lgot
SEQ EQ	2	574	574	574	574	574	574

PDB annotation	SUBUNIT; COMPLEX (GTP- BINDING/TRANSDUCER), G PROTEIN, HETEROTRIMER 2 SIGNAL TRANSDUCTION	HYDROLASE PROLYL ENDOPEPTIDASE, POST-PROLINE CLEAVING PROLYL OLIGOPEPTIDASE, AMNESIA, ALPHAMBETA-HYDROLASE, BETA- 2 PROPELLER	OXIDOREDUCTASE ENZYME, NITRITE REDUCTASE, OXIDOREDUCTASE, DENITRIFICATION, 2 ELECTRON TRANSPORT, PERIPLASMIC		TRANSMEMBRANE PROTEIN COLICIN, BACTERIOCIN, ION CHANNEL FORMATION, TRANSMEMBRANE 2 PROTEIN				INSECT IMMUNITY INSECT IMMUNITY, LPS-BINDING, HOMOPHILIC ADHESION	INSECT IMMUNITY INSECT IMMUNITY, LPS-BINDING, HOMOPHILIC ADHESION	RECEPTOR RECEPTOR, SIGNAL	CYTOKINES, THIRD 2 N-TERMINAL	DOMAIN, TRANSMEMBRANE, GLYCOPROTEIN	T-CELL SURFACE GLYCOPROTEIN IMMUNOGLOBULIN FOLD,
Coumpound		PROLYL OLIGOPEPTIDASE; CHAIN: A;	CYTOCHROME CDI NITRITE REDUCTASE; CHAIN: A, B;	٠	COLICIN IA; CHAIN: NULL;	DEFENSIN DEFENSIN /HNP\$-31DFN 3	DEFENSIN DEFENSIN /HNP\$-3 1DFN 3	DEFENSIN DEFENSIN /HNP\$-3 IDFN 3	HEMOLIN; CHAIN: A, B;	HEMOLIN; CHAIN: A, B;	GP130; CHAIN: NULL;			T-CELL SURFACE GLYCOPROTEIN CD4;
SeqFold score					98.21	59.92			111.17					
PMF	T.	0.05	-0.19				-	-		0.05	-0.08			-0.01
Verify score		-0.33	0.43				-0.35	-0.35		0	0.22			0.43
PSI- BLAST		0.00056	5.60E-79		1.70E-10	1.10E-12	1.10E-12	5.10E-11	1.70E-47	1.70E-47	5.60E-11			1.40E-19
End		172	478		648	101	101	101	435	416	434			220
Start AA		=	104		49	72	73	73	30	32	344			49
Chain		V V	٧			<	V	٧	٧	∢				
PDB ID		1qfm	1 qks		lcii	1dfn	ugp1	1dfn	1bih	15ih	1bj8			1cdy
SEQ B		574	574		575	577	217	577	578	578	278			578

. PDB annotation	TRANSMEMBRANE, GLYCOPROTEIN, T-CELL, 2 MHC, LIPOPROTEIN, T- CELL SURFACE GLYCOPROTEIN		CELL ADHESION NEURAL CELL ADHESION	CELL ADHESION NEURAL CELL ADHESION	IMMUNE SYSTEM ABZYME TRANSITION STATE ANALOG, IMMUNE SYSTEM	GROWTH FACTOR/GROWTH FACTOR RECEPTOR FGF, FGFR, IMMUNOGLOBULIN-LIKE, SIGNAL TRANSDUCTION, 2 DIMERIZATION, GROWTH FACTOR/GROWTH FACTOR RECEPTOR	GROWTH FACTOR/GROWTH FACTOR RECEPTOR FGF, FGFR, IMMUNOGLOBULIN-LIKE, SIGNAL TRANDUCTION, 2 DIMERIZATION, GROWTH FACTOR/GROWTH FACTOR	GROWTH FACTOR/GROWTH FACTOR RECEPTOR FGF, FGFR, IMMUNOGLOBULIN-LIKE, SIGNAL TRANSDUCTION, 2 DIMERIZATION, GROWTH FACTOR/GROWTH FACTOR RECEPTOR	CELL ADHESION NCAM; NCAM, IMMUNOGLOBULIN FOLD, GLYCOPROTEIN
Coumpound	CHAIN: NULL;	NEURAL ADHESION MOLECULE DROSOPHILA NEUROGLIAN (CHYMOTRYPTIC FRAGMENT CONTAINING THE ICFB 3 TWO AMINO PROXIMAL FIBRONECTIN TYPE III REPEATS ICFB 4 (RESIDUES 610 - 814)) ICFB 5	AXONIN-1; CHAIN: A;	AXONIN-1; CHAIN: A;	7C8 FAB FRAGMENT; SHORT CHAIN; CHAIN: A, C; 7C8 FAB FRAGMENT; LONG CHAIN; CHAIN: B, D	FIBROBLAST GROWTH FACTOR 2; CHAIN: A, B; FIBROBLAST GROWTH FACTOR RECEPTOR 1; CHAIN: C, D;	FIBROBLAST GROWTH FACTOR 2; CHAMI: A, B; FIBROBLAST GROWTH FACTOR RECEPTOR 1; CHAIN: C, D;	FIBROBLAST GROWTH FACTOR 2; CHAIN: A, B; FIBROBLAST GROWTH FACTOR RECEPTOR 1; CHAIN: C, D;	NEURAL CELL ADHESION MOLECULE; CHAIN: A, B, C, D;
SeqFold score							·		
PMF		0.39	-0.11	0	-0.15	-1.41	0.43	0.34	0.99
Verify score		0.25	60.0	90.0	0.11	0.1	0.18	0.1	0.16
PSI- BLAST		7.00E-20	1.40E-48	1.40E-50	6.80E-13	1.40E-44	3.405-28	1.40E-30	5.60E-30
End		435	522	436	328	340	225	220	214
Start AA		252	137	32	140	134	31	36	33
Chain			A	Ą	<	ပ	Ω	Ω	∢ .
PDB		ਜੂਰ ਜੂਰ	1cs6	les6	1ct8	lcvs	lcvs	1cvs	lepf
SEQ D		578	578	578	578	578	578	578	578

PDB annotation	GROWTH FACTOR/GROWTH FACTOR RECEPTOR FGF2; FGFR2; MMUNOGLOBULIN (GG)LIKE DOMAINS BELONGING TO THE I-SET 2 SUBGROUP WITHIN IG-LIKE DOMAINS, B-TREFOIL FOLD	GROWTH FACTOR/GROWTH FACTOR RECEPTOR FGF2; FGFR2; INMUNOGLOBULIN (IG)LIKE DOMAINS BELONGING TO THE I-SET 2 SUBGROUP WITHIN IG-LIKE DOMAINS, B-TREFOIL FOLD	GROWTH FACTOR/GROWTH FACTOR RECEPTOR FGF!; FGFR!; IMMUNOGLOBULIN (IG) LIKE DOMAINS BELONGING TO THE I-SET 2 SUBGROUP WITHIN IG-LIKE DOMAINS, B-TREFOIL FOLD	HORMONE/GROWTH FACTOR/HORMONE RECEPTOR 4- HELICAL BUNDLE, ALPHA HELICAL BUNDLE, TERNARY COMPLEX, FN 2 III DOMAINS, BETA SHEET DOWAINS, CYTOKINE-RECEPTOR COMPLEX	CONTRACTILE PROTEIN IMMUNOGLOBULIN FOLD, BETA BARREL	CELL ADHESION PROTEIN RGD, EXTRACELLULAR MATRIX IFNF 18	HEPARIN AND INTEGRIN BINDING HEPARIN AND INTEGRIN BINDING	-			IMMUNOGLOBULIN INTACT
Coumpound	FIBROBLAST GROWTH FACTOR 2; CHAIN: A, B, C, D; FIBROBLAST GROWTH FACTOR RECEPTOR 2; CHAIN: E, F, G, H;	FIBROBLAST GROWTH FACTOR 2; CHAIN: A, B, C, D; FIBROBLAST GROWTH FACTOR RECEPTOR 2; CHAIN: E, F, G, H;	FIBROBLAST GROWTH FACTOR 1; CHAIN: A, B; FIBROBLAST GROWTH FACTOR RECEPTOR 1; CHAIN: C, D;	PLACENTAL LACTOGEN; CHAIN: A; PROLACTIN RECEPTOR: CHAIN: B, C;	TELOKIN; CHAIN: A	FIBRONECTIN; IFNF 6 CHAIN: NULL; IFNF 7	FIBRONECTIN; CHAIN: A;	T LYMPHOCYTE ADHESION GLYCOPROTEIN CD2 (HUMAN) 1HNF 3	T LYMPHOCYTE ADHESION GLYCOPROTEIN CD2 (RAT) IFING 3	T LYMPHOCYTE ADHESION GLYCOPROTEIN CD2 (RAT) IHNG 3	IGG1 INTACT ANTIBODY
SeqFold score											95.64
PMF	0.25	0.48	0.01	0.17	-0.05	0.46	0.04	0.29	-0.12	0.15	
Verify score	60:0	-0.06	0.03	0.29	90.0	0.24	0.24	0.3	0.05	0.15	
PSI- BLAST	4.20E-30	2.80E-31	5.60E-28	1,40E-11	6.80E-18	5.60E-15	1.40E-12	1.40E-18	4.20E-14	2.80E-24	3.40E-10
End	220	228	220	436	131	432	432	211	292	227	454
Start	38	36	29	267	27	264	264	47	143	49	34
Chain ID	ய	ق ق	ပ	æ	4		٧		⋖	4	В
PDB ID	lev2	lev2	levt	1f6f	1fhg	1fnf	1fnb	1haf	1hng	1hng	ligy
SEQ S	578	578	578	578	578	578	878	878	578	578	578

PDB annotation	IMMUNOGLOBULIN, V REGION, C REGION, HINGE REGION	COMPLEX (IMMUNOGLOBULIN/RECEPTOR) IMMUNOGLOBULIN FOLD, IRANSMEMBRANE, GLYCOPROTEIN, RECEPTOR, 2 SIGNAL, COMPLEX (IMMUNOGLOBULIN/RECEPTOR)		CELL ADHESION PROTEIN CELL ADHESION PROTEIN, RGD, EXTRACELLULAR MATRIX, 2 HEPARIN-BINDING, GLYCOPROTEIN	MUSCLE PROTEIN CONNECTIN, NEXTMS; CELL ADHESION, LYCOPROTEIN, TRANSMEMBRANE, REPEAT, BRAIN, 2 INMUNOGLOBULIN FOLD, ALTERNATIVE SPLICING, SIGNAL, 3 MUSCLE PROTEIN	MUSCLE PROTEIN CONNECTIN, NEXTMS; CELL ADHESION, LYCOPROTEIN, TRANSMEMBRANE, REPEAT, BRAIN, 2 IMMUNOGLOBULIN FOLD, ALTERNATIVE SPLICING, SIGNAL, 3 MUSCLE PROTEIN	MUSCLE PROTEIN CONNECTIN, NEXTMS; CELL ADHESION, LYCOPROTEIN, TRANSMEMBRANE, REPEAT, BRAIN, 2 IMMUNOGLOBULIN FOLD, ALTERNATIVE SPLICING, SIGNAL, 3 MUSCLE PROTEIN	STRUCTURAL PROTEIN INTEGRIN, HEMIDESMOSOME, FIBRONECTIN, CARCINOMA, STRUCTURAL 2 PROTEIN
	IMMUN	COMPLEX (IMMUNO IMMUNOC TRANSME RECEPTOI		CELL / ADHES EXTRA HEPAR	MUSCI NEXTR GLYCC REPEA IMMUI ALTER MUSCI	MUSC BLYCO GLYCO REPEA IMMU ALTER MUSC		
Coumpound	MAB61.1.3; CHAIN: A, B, C, D	INTERLEUKIN-1 BETA; CHAIN: A; TYPE 1 INTERLEUKIN-1 RECEPTOR; CHAIN: B;	IMMUNOGLOBULIN IMMUNOGLOBULIN G1 (IGG1) (MCG) WITH A HINGE DELETION 1MCO 3	FIBRONECTIN; CHAIN: NULL;	TITIN; CHAIN: NULL;	TITIN; CHAIN: NULL;	TITIN; CHAIN: NULL;	INTEGRIN BETA-4 SUBUNIT; CHAIN: A, B;
SeqFold score			96.36					
PMF		-0.12	i	0.33	88.0	-0.06	-0.13	0.52
Verify score		0.03		0.39	0.29	90.00	0.13	0.3
PSI- BLAST		6.80E-34	3.40E-10	1.30E-17	1.40E-18	4.20E-15	6.80E-15	1.40E-20
End		338	423	432	227	132	132	450
Start		22	32	264	139	31 .	31	251
Chain ID		B	Н					∢
PDB DD		1itb	1mc o	1mfn	Inct	Inct	Inct	1983
SEQ	Ö	578	578	578	578	578	578	578

									
PDB annotation	STRUCTURAL PROTEIN TENASCIN, FIBRONECTIN TYPE-III, HEPARIN, EXTRACELLULAR 2 MATRIX, ADHESION, FUSION PROTEIN, STRUCTURAL PROTEIN				GLYCOPROTEIN CD4: IMMUNOCLOBULIN FOLD, TRANSMEMBRANE, GLYCOPROTEIN, T-CELL, 2 MHC LIPOPROTEIN, POLYMORPHISM	MUSCLE PROTEIN IMMUNOGLOBULIN SUPERFAMILY, I SET, MUSCLE PROTEIN	MUSCLE PROTEIN INMUNOGLOBULIN SUPERFAMILY, I SET, MUSCLE PROTEIN	NERVE GROWTH FACTOR/TRKA COMPLEX, TRKA RECEPTOR, NERVE GROWTH FACTOR, CYSTEINE KNOT, 2 IMMUNOGLOBULIN LIKE DOMAIN, NERVE GROWTH FACTOR/TRKA COMPLEX	CELL ADHESION ICAM-2; IMMUNOGLOBULIN FOLD, CELL ADHESION, GLYCOPROTEIN, 2 TRANSMEMBRANE, REPEAT, SIGNAL
Coumpound	TENASCIN; CHAIN: A, B;	MUSCLE PROTEIN TITIN MODULE MS (CONNECTIN) ITNM 3 (NMR, MINIMIZED AVERAGE STRUCTURE) ITNM 4 ITNM 58	MUSCLE PROTEIN TITIN MODULE MS (CONNECTIN) ITIMA 3 (WRR, MINIMIZED AVERAGE STRUCTURE) ITIMA 4 ITIM 58	GLYCOPROTEIN FIBRONECTIN (TENTH TYPE III MODULE) (NMR, 36 STRUCTURES) 1TTF 3	T-CELL SURFACE GLYCOPROTEIN CD4; CHAIN: A, B;	TWITCHIN 18TH IGSF MODULE; CHAIN: NULL;	TWITCHIN 18TH IGSF MODULE; CHAIN: NULL;	NERVE GROWTH FACTOR; CHAIN: V, W; TRKA RECEPTOR; CHAIN: X, Y;	INTERCELLULAR ADHESION MOLECULE-2; CHAIN: NULL;
SeqFold score									
PMF	0.82	0.89	0.03	0.05	-0.12	0.78	0.1	-0.01	0.23
Verify score	0.52	0.39	-0.04	0.2	0.04	0.52	0.23	0.02	0.18
PSI- BLAST	2.80E-17	7.00E-18	8.40E-15	1.40E-10	2.80E-20	2.80E-17	2.80E-15	5.60E-18	1.10E-30
End	435	722	132	434	281	226	132	228	228
Start AA	264	139	33	346		139	31		31
Chain ID	. ∢		,		V			×	
PDB ID	1914	1tnm	Ithm	Ħ	lwio	lwit	lwit	w w	lzxq
SEQ	578	578	578	578	578	578	578	578	578

PDB annotation	IMMUNE SYSTEM PS8 NATURAL KILLER CELL RECEPTOR, KIR, NATURAL KILLER RECEPTOR, INHIBITORY RECEPTOR, 2 IMMUNOGLOBULIN	IMMUNE SYSTEM CD32; RECEPTOR, FC, CD32, IMMUNE SYSTEM	PROTEIN BINDING ED-B, FIBRONECTIN, TYPEIII DOMAIN, ANGIOGENESIS, PROTEIN 2 BINDING	COAGULATION FACTOR	CELL ADHESION NCAM DOMAIN 1; CELL ADHESION, GLYCOPROTEIN, HEPARIN-BINDING, GPI-ANCHOR, 2 NEURAL ADHESION MOLECULE, IMMUNOGLOBULIN FOLD, SIGNAL		CELL ADHESION PROTEIN NCAM MODULE 2; CELL ADHESION, GLYCOPROTEIN, HEPARIN-BINDING, GPI-ANCHOR, 2 NEURAL ADHESION MOLECULE, IMMUNOGLOBULIN FOLD, HOMOPHILIC 3 BINDING, CELL ADHESION PROTEIN	CELL ADHESION PROTEIN NCAM MODULE 2; CELL ADHESION, GLYCOPROTEIN, HEPARIN-BINDING, GPLYANCHOK, 2 NEURAL ADHESION MOLECULE, IMMUNOGLOBULIN FOLD, HOMOPHILIC 3 BINDING, CELL ADHESION PROTEIN	
Coumpound	MHC CLASS I NK CELL RECEPTOR PRECURSOR; CHAIN: A;	FC GAMMA RIB; CHAIN: A;	FIBRONECTIN; CHAIN: A;	HUMAN TISSUE FACTOR; 2HFT 4 CHAIN: NULL; 2HFT 5	NEURAL CELL ADHESION MOLECULE; CHAIN: NULL;	HORMONERECEPTOR HUMAN GROWFLEXED HORMONE COMPLEXED WITH ITS RECEPTOR 3HHR 3 (EXTRACELLULAR DOMAIN) 3HHR 4	NEURAL CELL ADHESION MOLECULE, LARGE ISOFORM; CHAIN: A;	NEURAL CELL ADHESION MOLECULE, LARGE ISOFORM; CHAIN: A;	VIRUS EQUINE HERPES VIRUS-1 (C3HC4, OR RING DOMAIN) ICHC 3 (NMR, 1
SeqFold score									
PMF	-0.02	99.0	-0.12	-0.08	0.98	-0.09	0.31	0.06	0.01
Verify score	0.25	0.1	0.45	0.13	0.64	90.00	0.06	0.04	-0.56
PSI- BLAST	2.80E-25	2.80E-33	1.40E-12	2.80E-12	5.60E-18	5.60E-17	4.20E-18	7.00E-17	1.00E-09
End	220	228	435	436	227	436	220	134	103
Start AA	30	31	343	264	139			32	19
Chain	∢	¥	Ą			Ø	∢	∢	
PDB TD	2dli	2fcb	2fnb	2hft	Znc m	3hhr	3nc m	3nc B	1chc
SEQ S B S	578	578	578	578	578	578	878	578	579

PDB annotation		TRANSCRIPTION FACTOR BTF2 P44 SUBUNIT; BASIC TRANSCRIPTION FACTOR, ZINC BINDING PROTEIN	MANA BININIS PROTEIN VIDI	RECOMBINATION ACTIVATING	PROTEIN I: RAG1, V(D)J	RECOMBINATION, ANTIBODY, MAD,	RING FINGER, 2 ZINC BINUCLEAR	CLISTER ZINC FINGER DNA-	BINDING PROTEIN	DNA-BINDING PROTEIN V(D)J	RECOMBINATION ACTIVATING	PROTEIN 1; RAG1, V(D)J	RECOMBINATION, ANTIBODY, MAD,	RING FINGER, 2 ZINC BINUCLEAR	CLUSTER, ZINC FINGER, DNA-	BINDING PROTEIN	DNA-BINDING PROTEIN V(D)J	RECOMBINATION ACTIVATING	PROTEIN 1; RAG1, V(D)J	RECOMBINATION, ANTIBODY, MAD,	RING FINGER, 2 ZINC BINUCLEAR	CLUSTER, ZINC FINGER, DNA-	BINDING PROTEIN		CALCIUM-BINDING PROTEIN CALB;	CALCIOM++/FROSTHOLIFID BINDING	PROTEIN	ENDOCYTOSIS/EXOCYTOSIS	SYNAPTOTAGMIN, C2-DOMAIN,	EXOCYTOSIS, NEUROTRANSMITTER	2 RELEASE,	ENDOCYTOSIS/EXOCYTOSIS	⊢	_	HYDROLASE	TRANSFERASE CALCIUM++,	PHOSPHOLIPID BINDING PROTEIN, CALCIUM-BINDING 2 PROTEIN,
Coumpound	STRUCTURE) ICHC 4	TFIIH P44 SUBUNIT; CHAIN: A;	DACT. CHAINLANT I.	KAGI; CHAIN: NOLL;						RAG1; CHAIN: NULL;							RAG1; CHAIN: NULL;					,			PROTEIN KINASE C (BETA);	CHAIN: A, B;		SYNAPTOTAGMIN I: CHAIN:	A:				CYTOSOLIC	PHOSPHOLIPASE A2; CHAIN:	A, B;	PROTEIN KINASE C, ALPHA	TYPE; CHAIN: A;
SeqFold score																									 												
PMF		0.19	5	0.33						0.29							0.13							-	0.94			0.50	7				0.13			86.0	İ
Verify score		0		- - - - -						-0.78							0.01								0.83			900	3				0.05			19.0	
PSI- BLAST		0.00051		0.0037						0.0007							3.40E-07								1.70E-30			1 70E 23					1.70E-15			3.40E-31	
End		102		268						92						,	128								1443			1442	7				1444			1443	
Start		51		223						59							63								1329			1325	77.				1342			1329	
Chain ID		¥																							٧			\ \	ζ				4			Ą	
PDB TD		1.00 E+53		lrmd						lrmd							1md								1825			1,4,7	1601				Icjy			1dsy	
SEQ B B		879		579						579							579		_						280			Ş	8				280			280	

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PDB annotation	PHOSPHATIDYLSERINE, PROTEIN KINASE C	PHOSPHONOSITIDE 3-KINASE GAMMA PTDINS-3-KINASE P110, P13K, P13K; PHOSPHOINOSITIDE 3-KINASE GAMMA, SECONDARY MESSENGER 2 GENERATION, P13K, P1 3K, WORTMANNIN	PHOSPHOINOSITIDE 3-KINASE GAMMA PTDINS-3-KINASE P110, P13K; PHOSPHOINOSITIDE 3-KINASE GAMMA, SECONDARY MESSENGER 2 GENERATION, P13K, P13K	HYDROLASE CALB DOMAIN; HYDROLASE, C2 DOMAIN, CALB DOMAIN		ENDOCYTOSIS/EXOCYTOSIS C2- DOMAINS, C2B-DOMAIN, RABPHILIN, ENDOCYTOSIS/EXOCYTOSIS	SIGNALING PROTEIN GUANINE NUCLEOTIDE- BINDING PROTEIN 1; GBP, GTP HYDROLYSIS, GDP, GMP, INTERFERON INDUCED, DYNAMIN 2 RELATED, LARGE GTPASE FAMILY, SIGNALING PROTEIN	SIGNALING PROTEIN GBP, GTP HYDROL YSIS, GDP, GMP, INTERFERON INDUCED, DYNAMIN 2 RELATED, LARGE GTPASE FAMIL Y. GMPPNP, GPPNHP.	GOT GURGIST GAROTTE	PHOSPHOTRANSFERASE PHOSPHOGLUCOMUTASE; 3PMG 6 PHOSPHOGLUCOMITASE; 3PMG 13	PHOSPHOTRANSFERASE PHOSPHOGLUCOMUTASE; 3PMG 6
Coumpound		PHOSPHATIDYLINOSITOL 3- KINASE CATALYTIC SUBUNIT; CHAIN: A;	PHOSPHATIDYL,NOSITOL 3- KINASE CATALYTIC SUBUNIT; CHAIN: A;	PHOSPHOLIPASE A2; CHAIN: NULL;	CALCIUM/PHOSPHOLIPID BINDING PROTEIN SYNAPTOTAGMIN I (FIRST C2 DOMAIN) (CALB) 1RSY 3	RABPHILIN 3-A; CHAIN: A;	INTERFERON-INDUCED GUANYLATE-BINDING PROTEIN I; CHAIN: A;	INTERFERON-INDUCED GUANYLATE-BINDING PROTEIN 1; CHAIN: A;	,	ALPHA-D-GLUCOSE-1,6- BISPHOSPHATE; 3PMG 4 CHAIN: A. B: 3PMG 5	ALPHA-D-GLUCOSE-1,6- BISPHOSPHATE; 3PMG 4
SeqFold score											129.26
PMF		1202.08	-1.41	0.64	0.59	0.39	-0.2	-0.2	,	7.0-	
Verify score		0.15	0.15	-0.02	0.06	0.1	0.27	0.47	3	67:0	
PSI- BLAST		0	0	1.70E-15	8.50E-24	5.10E-29	0	0		-	0
End AA		1175	1175	1444	1442	1441	577	577	707	Š	611
Start AA		214		1342	1321	1328	9	7	y	₽	53
Chain ID		¥	4			٧	∢	4	4	٤	A
PDB ID		le7u	le8y	Iriw	Irsy	3грь	1dg3	<u>ν</u>	3nm	8	3pm g
SEQ D NO:		280	580	280	280	280	2885	585	486	3	586

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PDB annotation	PHOSPHOGLUCOMUTASE 3PMG 13	TRANSCRIPTION REGULATION	PROTO-ONCOGENE, NUCLEAR BODIES (PODS), LEUKEMIA, 2 TRANSCRIPTION REGULATION	TRANSCRIPTION REGULATION PROTO-ONCOGENE, NUCLEAR BODIES (PODS), LEUKEMIA, 2 TRANSCRIPTION REGULATION		LIGASE CBL, UBCH7, ZAP-70, E2, UBIQUITIN, E3, PHOSPHORYLATION, 2 TYROSINE KINASE, UBIQUITINATION, PROTEIN DEGRADATION,	METAL BINDING PROTEIN RING FINGER PROTEIN MATI; RING FINGER (C3HC4)	METAL BINDING PROTEIN MING FINGER PROTEIN MATI; RING FINGER (C3HC4)	LIM DOMAIN CONTAINING PROTEINS LIM DOMAIN CONTAINING PROTEINS, METAL-BINDING PROTEIN, ZINC 2 FINGER	LIM DOMAIN CONTAINING PROTEINS LIM DOMAIN CONTAINING PROTEINS, METAL-BINDING PROTEIN, ZINC 2 FINGER	CONTRACTILE LIM DOMAIN, CRP, NMR, MUSCLE DIFFERENTIATION, CONTRACTILE	METAL-BINDING PROTEIN LIM DOMAIN CONTAINING PROTEINS 1CTL 15
Coumpound	CHAIN: A, B; 3PMG 5	TE ANSCEIPTION FACTOR	PML; CHAIN: NULL;	TRANSCRIPTION FACTOR PML; CHAIN: NULL;	VIRUS EQUINE HERPES VIRUS-1 (C3HC4, OR RING DOMAIN) ICHC 3 (NMR, 1 STRUCTURE) ICHC 4	SIGNAL TRANSDUCTION PROTEIN CBL: CHAIN: A: ZAP-70 PEPTIDE: CHAIN: B; UBIQUITIN-CONJUGATING ENZYME E12-18 KDA 11BCH7: CHAIN: C:	CDK-ACTIVATING KINASE ASSEMBLY FACTOR MATI; CHAIN: A;	CDK-ACTIVATING KINASE ASSEMBLY FACTOR MATI; CHAIN: A;	QCRP2 (LIM1); CHAIN: NULL;	QCRP2 (LIM1); CHAIN: NULL;	CRP1; CHAIN: A;	AVIAN CYSTEINE RICH PROTEIN; ICTL 3
SeqFold score				,								
PMF			570	0.11	0.98	6.0	0.48	0.09	0.62	0	0.18	0.16
Verify score			-0.73	-0.62	0.24	0.25	0.19	-0.44	-0.16	, 0.1	-0.11	-0.11
PSI- BLAST			8.50E-06	8.40E-08	8.50E-15	1.705-10	8.40E-08	1.20E-05	9.80E-17	1.40E-11	4.20E-11	1.10E-12
End			65	56	. 09	99	2	72	292	662	659	629
Start AA			16		16	<u>8</u>	17	. 18	536	597	597	969
Chain						₹	<	A			<	
PDB U	\top	T	1bor	1bor	1chc	1fbv	1g25	1825	1a7i	la7i	168t	1ct
SEQ	ÿ		587	587	587	587	587	587	292	292	592	292

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PDB annotation	SIGNALING PROTEIN LIM DOMAIN CONTAINING PROTEINS, METAL- BINDING PROTEIN	SIGNALING PROTEIN LIM DOMAIN CONTAINING PROTEINS, METAL- BINDING PROTEIN	METAL-BINDING PROTEIN CRIP; METAL-BINDING PROTEIN, LIM DOMAIN PROTEIN	METAL-BINDING PROTEIN CRIP; METAL-BINDING PROTEIN, LIM DOMAIN PROTEIN	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)		ANTI-ONCOGENE CELL CYCLE, ANTI- ONCOGENE, REPEAT, ANK REPEAT	COMPLEX (TRANSCRIPTION REGULATION/DNA) GABPALPHA; GABPBETA I: COMPLEX	(IKANSCKIPTION REGULATIONDNA), DNA-BINDING, 2 NUCLEAR PROTEIN, ETS DOMAIN, ANK YRIN REPEATS. TRANSCRIPTION	3 FACTOR	TUMOR SUPPRESSOR TUMOR SUPPRESSOR, CDK4/6 INHIBITOR, ANKYRIN MOTIF	COMPLEX (KINASE/ANTI- ONCOGENE) CDK6; P16INK44, MTS1; CYCLIN DPERDIENT KINASE,	CICLIN DEFENDENT ANASSE INHIBITORY 2 PROTEIN, CDK, INK4, CELT CYCLE ACIT TIPLE TIMOR	SUPPRESSOR, 3 MTS1, COMPLEX (KINASE/ANTI-ONCOGENE) HEADER	COMPLEX (INHIBITOR PROTEINIKINASE) INHIBITOR
Coumpound	CYSTEINE AND GLYCINE- RICH PROTEIN CRP2; CHAIN: A:	CYSTEINE AND GLYCINE- RICH PROTEIN CRP2; CHAIN: A;	CYSTEINE RICH INTESTINAL PROTEIN; CHAIN: NULL;	CYSTEINE RICH INTESTINAL PROTEIN; CHAIN: NULL;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;		TUMOR SUPPRESSOR P16INK4A; CHAIN: NULL;	GA BINDING PROTEIN ALPHA; CHAIN: A; GA BINDING PROTEIN BETA 1;	CHAIN: B; DNA; CHAIN: D, E;		P19INK4D CDK4/6 INHIBITOR; CHAIN: NULL;	CYCLIN-DEPENDENT KINASE 6; CHAIN: A; MULTIPLE TUMOR	SUPPRESSOR; CHAIN: B;		CYCLIN-DEPENDENT KINASE 6; CHAIN: A;
SeqFold score															
PMF	0.8	0.23	0.29	0.07	-0.19		0.72	0.1			0.76	0.82			0.84
Verify score	0.43	-0.27	-0.42	0.15	0.16		1.0	0.35			0.28	0.02			0.23
PSI- BLAST	1.40E-16	1.10E-13	1.30E-19	4.20E-13	1.20E-10		0.00014	0.00014			0.00011	9.80E-05	<u>-</u>		0.00011
End	592	199	609	663	395		370	369			370	367			367
Start	535	595	537	296	367		315	315			315	315			315
Chain	<	4			Ð			8				æ			æ
PDB	1cx	1cxx	Lii.	limi	1me y		1a5e	lawc			1bd8	1bi7			1blx
SEQ	292 292	592	592	592	592		593	593			593	593			293

PDB annotation	PROTEIN, CYCLIN-DEPENDENT KINASE, CELL CYCLE 2 CONTROL, ALPHA/BETA, COMPLEX (INHIBITOR PROTEIN/KINASE)	COMPLEX (INHIBITOR PROTEINKINASE) INHIBITOR PROTEIN/KINASE) INHIBITOR PROTEIN, CYCLIN-DEPENDENT KINASE, CELL CYCLE 2 CONTROL, ALPHA/BETA, COMPLEX (INHIBITOR PROTEIN/KINASE)	SIGNALING PROTEIN HELIX-TURN- HELIX, ANKYRIN REPEAT	METAL BINDING PROTEIN ZINC- BINDING MODULE, ANKYRIN REPEATS, METAL BINDING PROTEIN	TRANSCRIPTION FACTOR P65; P50D; TRANSCRIPTION FACTOR, IKBNFKB COMPLEX	COMPLEX (TRANSCRIPTION REGIANK REPEAT) COMPLEX (TRANSCRIPTION REGULATION/ANK REPEAT), ANKYRIN 2 REPEAT HELIX	KINASE KINASE, SIGNAL TRANSDUCTION, CALCIUM/CALMODULIN	
Coumpound	P19NK4D; CHAIN: B;	CYCLIN-DEPENDENT KINASE 6; CHAIN: A; PI9INK4D; CHAIN: B;	CYCLIN-DEPENDENT KINASE 4 INHIBITOR B; CHAIN: A;	PYK2-ASSOCIATED PROTEIN BETA; CHAIN: A;	NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF-KAPPA-B P50D SUBUNIT; CHAIN: C; I- KAPPA-B-ALPHA; CHAIN: D;	NP-KAPPA-B P65; CHAIN: A, C; NF-KAPPA-B P50; CHAIN: B, D; I-KAPPA-B-ALPHA; CHAIN: E, F;	CALCIUM/CALMODULIN- DEPENDENT PROTEIN KINASE; CHAIN: NULL;	TRANSFERASE(PHOSPHOTR ANGFERASE) \$C-/AMP\$- DEPENDENT PROTEIN KINASE (E.C.2.7.1.37) (\$CATALYTIC SUBUNIT) ALPHA ISOENZYME MUTANT WITH SER 139 IAPM 4 REPLACED BY ALA (\$1394\$) COMPLEX WITH THE PEPTIDE 1APM 5 INHIBITOR PKI(\$-24) AND THE DETERGENT MEGA-8
SeqFold score		·						
PMF		0.76	0.21	97.0	0.98	0.59	_	_
Verify score		0.42	-0.02	0.4	0.31	0.25	0.15	0.23
PSI- BLAST		0.00011	1.40E-05	8.40E-05	0.00014	2.80E-06	5.10E-90	
End		370	370	370	370	381	302	318
Start		315	315	315	315	315	19	25
Chain ID		В	¥	4	Ω	a		ω .
PDB		×J91	1d9s	1dcq	1.jkm	lufi	1a06	lap m .
SEQ.	Ö	593	593	593	593	593	594	594

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PDB annotation			PROTEIN KINASE CDK2; PROTEIN KINASE, CELL CYCLE, PHOSPHORYLATION, STAUROSPORUE, 2 CELL DIVISION, MITOSIS, INHIBITION	COMPLEX (INHIBITOR PROTEIN/KINASE) INHIBITOR PROTEIN, CYCLIN-DEPENDEN KINASE, CELL CYCLE 2 CONTROL, ALPHABETA, COMPLEX (INHIBITOR PROTEIN/KINASE)			
Coumpound	1APM 6	TRANSFERASE(PHOSPHOTR ANSFERASE) \$C-\AMP\$- DEPENDENT PROTEIN KINASE (E.C.2.7.1.37) (\$CAPK\$) 1APM 3 (\$CAPK\$) 1APM 3 (CATALYTIC SUBUNIT) ALPHA ISOENZYME MUTANT WITH SER 139 1APM 4 REPLACED BY ALA (\$1394\$) COMPLEX WITH THE PEPTIDE 1APM 5 INHBITOR PKI(5-24) AND THE DETERGENT MEGA-8 1APM 6	CYCLIN-DEPENDENT PROTEIN KINASE 2; CHAIN: NULL;	CYCLIN-DEPENDENT KINASE 6; CHAIN: A; PI9INK4D; CHAIN: B;	PHOSPHOTRANSFERASE CAMP-DEPENDENT PROTEIN KINASE CATALYTIC SUBUNIT ICMK 3 (E.C.2.7.1.37) ICMK	PHOSPHOTRANSFERASE CAMP-DEPENDENT PROTEIN KINASE CATALYTIC SUBUNIT ICMK 3 (E.C.2.7.1.37) ICMK 4	TRANSFERASE(PHOSPHOTR ANSFERASE) CAMP- DEPENDENT PROTEIN KINASE (E.C.2.1.137) (CAPK) ICTP 3 (CATALYTIC
SeqFold score		125.65	120.88	127.11	127.4		127.61
PMF						-	
Verify						0.33	
PSI- BLAST			1.40E-60	7.00E-56	0	0	0
End		325	309	325	328	318	328
Start AA:		v	27	22	-	25	7
Chain		<u>.</u>		∢	凹	ш	ш
PDB CD		I ap	laqı	1bk	lcm k	k k	1cp
SEQ EQ	5	594	594	594	594	594	594

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PDB annotation			ENDOCYTOSIS/EXOCYTOSIS NSECI; PROTEIN-PROTEIN COMPLEX, MULTI-SUBUNIT	ENDOCYTOSIS/EXOCYTOSIS SYNAPTOTAGMIN ASSOCIATED 35 KDA PROTEIN, P35A, THREE HELIX BUNDLE	ENDOCYTOSIS/EXOCYTOSIS SYNAPTOTAGMIN ASSOCIATED 35 KDA PROTEIN, P35A, THREE HELIX BUNDLE	TRANSFERASE KINASE DOMAIN, AUTOINHIBITORY FRAGMENT, HOMODIMER	TRANSFERASE KINASE DOMAIN, AUTOINHIBITORY FRAGMENT, HOMODIMER	PHOSPHOTRANSFERASE FGFRIK, FIBROBLAST GROWTH FACTOR RECEPTOR I; TRANSFERASE, TYROSINE-PROTEIN KINASE, ATP-BINDING, 2 PHOSPHORYLATION, RECEPTOR, PHOSPHOTRANSFERASE	PHOSPHOTRANSFERASE FGFRIK, FIBROBLAST GROWTH FACTOR RECEPTOR 1; TRANSFERASE, TYROSINE-PROTEIN KINASE, ATP-BINDING, 2 PHOSPHORYLATION,
Coumpound	SUBUNIT) ICTP 4	TRANSFERASE(PHOSPHOTR ANSFERASE) CAMP- DEPENDENT PROTEIN KINASE (C.2.7.1.37) (CAPK) ICTP 3 (CATAL.YTIC SUBUNIT) ICTP 4	SYNTAXIN BINDING PROTEIN I; CHAIN: A; SYNTAXIN IA; CHAIN: B;	SYNTAXIN-1A; CHAIN: A, B, C;	SYNTAXIN-1A; CHAIN: A, B, C;	SERNE/THREONINE- PROTEIN KINASE PAK- ALPHA; CHAIN: A, B; SERINE/THREONINE- PROTEIN KINASE PAK- ALPHA; CHAIN: C, D;	SERINE/THREONINE- PROTEIN KINASE PAK- ALPHA; CHAIN: A, B; SERINE/THREONINE- PROTEIN KINASE PAK- ALPHA; CHAIN: C, D;	FGF RECEPTOR 1; CHAIN: A, B;	FGF RECEPTOR 1; CHAIN: A, B;
SeqFold score								125.79	130.09
PMF			-0.11	-0.19	-0.18	-	1202.08		
Verify score		0.25	0.06	0.1	0.16	0.5	0.43		
PSI- BLAST		0	4.20E-12	1.40E-08	2.80E-09	5.60E-86	1.20E-67	5.10E-34	1.20E-40
End		318	009		617	302	300	287	286
Start AA		25	442	452	486	14	4	50	11
Chain ID		m	В	<	<	U	U	∢	æ
PDB ID		lctp	1dn1	lez3	1ez3	1f3m	163m	1fgk	1gk
SEQ EQ		594	594	594	594	594	594	594	594

PDB annotation	RECEPTOR, PHOSPHOTRANSFERASE	PROTEIN KINASE CDK2; TRANSFERASE, SERINE/THREONINE PROTEIN KINASE, ATT-BINDING, 2 CELL CYCLE, CELL DIVISION, MITOSIS, PHOSPHORYLATION	PROTEIN KINASE CDK2; TRANSFERASE, SERINE/THREONINE PROTEIN KINASE, ATP-BINDING, 2 CELL CYCLE, CELL DIVISION, MITOSIS, PHOSPHORYLATION	KINASE KINASE, TWITCHIN, INTRASTERIC REGULATION	KINASE KINASE, TWITCHIN, INTRASTERIC REGULATION	TRANSFERASE MITOGEN ACTIVATED PROTEIN KINASE; TRANSFERASE, MAP KINASE, SERINETHREONINE-PROTEIN KINASE, 2 P38	KINASE RABBIT MUSCLE PHOSPHORYLASE KINASE; GLYCOGEN METABOLISM, TRANSFERASE, SERINBTHREONINE- PROTEIN, 2 KINASE, ATP-BINDING, CALMODULIN-BINDING	KINASE RABBIT MUSCLE PHOSPHORYLASE KINASE; GLYCOGEN METABOLISM, TRANSFERASE, SERINGTHREONINE- PROTEIN, 2 KINASE, ATP-BINDING, CALMODULIN-BINDING	TRANSFERASE MITOGEN ACTIVATED PROTEIN KINASE, MAP 2, ERK2; TRANSFERASE, SERINE/THREONINE-PROTEIN KINASE, MAP KINASE, 2 ERK2	יייייייייייייייייייייייייייייייייייייי	KINASE KINASE, SIGNAL TRANSDUCTION, CALCIUM/CALMODULIN
Coumpound		HUMAN CYCLIN- DEPENDENT KINASE 2; CHAIN: NULL;	HUMAN CYCLIN- DEPENDENT KINASE 2; CHAIN: NULL;	TWITCHIN; CHAIN: NULL;	TWITCHIN; CHAIN: A, B;	MAP KINASE P38; CHAIN: NULL;	PHOSPHORYLASE KINASE; CHAIN: NULL;	PHOSPHORYLASE KINASE; CHAIN: NULL;	EXTRACELLULAR REGULATED KINASE 2; CHAIN: NULL;		CALCIUM/CALMODULIN- DEPENDENT PROTEIN KINASE; CHAIN: NULL;
SegFold score			137.2			130.84	127.79	·	122.57		
PMF		1		_	1.		·	_			1202.08
Verify		0.48		0.5	0.48			0.53			0
PSI- BLAST		6.80E-62	6.80E-62	3.40E-71	1.70E-71	1.20E-51	6.80E-88	6.80E-88	8.50E-50		1.00E-66
End		284	312	285	284	356	285	282	325		438
Start		25	27	70	20	9	61	24	10	0	140
Chain					4						
PDB		1hcl	1hcl	Ikoa	1kob	1p38	1 phk	1 phk	3erk		1a06
SEQ	ö	594	594	594	594	594	594	594	594		296

PDB annotation	TRANSFERASE TRANSFERASE, SERINE/THREONINE-PROTEIN KINASE, CASEIN KINASE, 2 SER/THR KINASE		PROTEIN KINASE CDK2; PROTEIN KINASE, CELL CYCLE, PHOSPHORYLATION, STAUROSPORINE, 2 CELL DIVISION, MITOSIS, INHIBITION	COMPLEX (KINASE/INHIBITOR) CDK6; P19INK4D; CYCLIN DEPENDENT KINASE, CYCLIN DEPENDENT KINASE INHIBITORY 2 PROTEIN, CDK, INK4, CELL CYCLE, COMPLEX (KINASE/INHIBITOR) HEADER HELIX	COMPLEX (INHIBITOR PROTEIN/KINASE) INHIBITOR PROTEIN, CYCLIN-DEPENDENT KINASE, CELL CYCLE 2 CONTROL, ALPHABETA, COMPLEX (INHIBITOR PROTEIN/KINASE)	TRANSFERASE CSK; PROTEIN KINASE, C-TERMINAL SRC KINASE, PHOSPHORYLATION, 2 STAUROSPORINE, TRANSFERASE	PHOSPHOTRANSFERASE PROTEIN
Coumpound	PROTEIN KINASE CK2/ALPHA-SUBUNIT; CHAIN: NULL;	TRANSFERASE(PHOSPHOTR ANSFERASE) \$C-\AMP\$- DEPENDENT PROTEIN KINASE (E.C.2.7.1.37) (\$C\APK\$) 1APM 3 (CATALYTIC SUBUNIT) ALPHA ISOENZYME MUTANT WITH SER 139 1APM 4 REPLACED BY ALA (\$S1394\$) COMPLEX WITH THE PEPTIDE 1APM 5 ITHE PEPTIDE 1APM 5 ITHE PEPTIDE 1APM 5 ITHE DETERGENT MEGA-8 1APM 6	CYCLIN-DEPENDENT PROTEIN KINASE 2; CHAIN: NULL;	CYCLIN-DEPENDENT KINASE 6; CHAIN: A, C; CYCLIN-DEPENDENT KINASE INHIBITOR; CHAIN: B, D;	CYCLIN-DEPENDENT KINASE 6; CHAIN: A; P19NK4D; CHAIN: B;	C-TERMINAL SRC KINASE; CHAIN: A;	CASEIN KINASE I DELTA;
SeqFold score							
PMF	0.22	0.49	0.76	0.05	6.3	0.46	0.34
Verify score	-0.44	-0.03	0.03	-0.06	0.2	0.01	-0.08
PSI- BLAST	4.20E-07	3.40E-100	8.50E-49	8.50E-32	1.70E-35	3.40E-30	5.60E-09
End AA	351	434	399	397	398	344	351
Start AA	234		146	149	149	146	198
Chain ID		B		⋖	<	<	A
PDB CD	1a6o	m m	laqi	1518	1bk	1byg	1cki
SEQ NO DE	296	596	596	596	596	965	596

PDB annotation	KINASE ICKI 18	TRANSFERASE STRESS-ACTIVATED PROTEIN KINASE-3, ERK6, ERK5; P38-GAMMA, GAMMA, PHOSPHORYLATION, MAP KINASE			TRANSFERASE KINASE DOMAIN, AUTOINHIBITORY FRAGMENT, HOMODIMER	PHOSPHOTRANSFERASE FGFR1K, FIBROBLAST GROWTH FACTOR FECEPTOR 1: TRANSFERASE, TYROSINE-PROTEIN KINASE, ATP-BINDING, 2 PHOSPHORYLATION, RECEPTOR, PHOSPHOTRANSFERASE	PHOSPHOTRANSFERASE FGFR IK, FIBROBLAST GROWTH FACTOR RECEPTOR 1; TRANSFERASE, TYROSINE-PROTEIN KINASE, ATP-BINDING, 2 PHOSPHOTRALATION, RECEPTOR, PHOSPHOTRANSFERASE		PROTEIN KINASE CDK2; TRANSFERASE, SERINE/THREONINE PROTEIN KINASE, ATP-BINDING, 2 CELL CYCLE, CELL DIVISION,
Coumpound	ICKI 6 CHAIN: A, B; ICKI 7	PHOSPHORYLATED MAP KINASE P38-GAMMA; CHAIN: A, B;	PHOSPHOTRANSFERASE CAMP-DEPENDENT PROTEIN KINASE CATALYTIC SUBUNIT ICMK 3 (E.C.2.7.1.37) ICMK 4	TRANSFERASE(PHOSPHOTR ANSFERASE) CAMP- DEPENDENT PROTEIN KINASE (E.C.2.7.137) (CAPK) ICTP 3 (CATALYTIC SUBUNIT) ICTP 4	SERINETHREONINE- PROTEIN KINASE PAK- ALPHA; CHAIN: A, B; SERINETHREONINE- PROTEIN KINASE PAK- ALPHA; CHAIN: C, D;	FGF RECEPTOR 1; CHAIN: A, B;	FGF RECEPTOR 1; CHAIN: A, B;	PROTO-ONCOGENE TYROSINE-PROTEIN KINASE ABL; CHAIN: A, B;	HUMAN CYCLIN- DEPENDENT KINASE 2; CHAIN: NULL;
SeqFold score							·		
PMF		0.09	0.37	0.48	0.89	0.22	0.33	0.51	0.7
Verify score		0.32	-0.11	0.18	0.35	0.14	-0.07	0.12	0.26
PSI- BLAST		1.50E-31	0	6.80E-97	6.80E-46	5.10E-31	7.00E-08	4.20E-06	1.70E-47
End		396	434	434	397	344	372	372	399
Start		163	122	122	148	147	506	192	146
Chain		A	ш	ш	ပ	В	B	4	
PDB DD	1	8 8 8	r Ich	1ctp	153m	1旗	1fgk	1fpu	lhcl
SEQ U	ä	596	969	965	296	296	965	296	296

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PDB annotation	MITOSIS, PHOSPHORYLATION	SERNE/THREONINE-PROTEIN KINASE CSBP, RK, P38; PROTEIN SER/THR-KINASE, SERINE/THREONINE-PROTEIN KINASE	TRANSFERASE INK3; TRANSFERASE, INK3 MAP KINASE, SERINE/THREONINE PROTEIN 2 KINASE.	TRANSFERASE JNK3; TRANSFERASE, JNK3 MAP KINASE, SERINE/THREONINE PROTEIN 2 KINASE	KINASE KINASE, TWITCHIN, INTRASTERIC REGULATION	KINASE KINASE, TWITCHIN, INTRASTERIC REGULATION	TRANSFERASE MITOGEN ACTIVATED PROTEIN KINASE; TRANSFERASE, MAP KINASE, SERINE/THREONINE-PROTEIN KINASE, 2 P38	KINASE RABBIT MÜSCLE PHOSPHORYLASE KINASE; GLYCOGEN METABOLISM, TRANSFERASE, SERINE/THIEONINE- PROTEIN, 2 KINASE, ATP-BINDING, CALMODILIN-BINDING	SERINE KINASE SERINE KINASE, TITIN, MUSCLE, AUTOINHIBITION	TRANSFERASE AAC; AMINOGLYCOSIDE 6'-N- ACETYLTRANSFERASE, ANTIBIOTIC 2 RESISTANCE, ACETYL COENZYME A	HYDROLASE ERA, GTPASE, RNA- BINDING, RAS-LIKE, HYDROLASE
Coumpound		P38 MAP KINASE; CHAIN: NULL;	C-JUN N-TERMINAL KINASE, CHAIN: NULL;	C-JUN N-TERMINAL KINASE; CHAIN: NULL;	TWITCHIN; CHAIN: NULL;	TWITCHIN; CHAIN: A, B;	MAP KINASE P38; CHAIN: NULL;	PHOSPHORYLASE KINASE; CHAIN: NULL;	TITIN; CHAIN: A, B;	AMINOGLYCOSIDE NG. ACETYLTRANSFERASE TYPE 1; CHAIN: A;	GTP-BINDING PROTEIN ERA; CHAIN: A, B;
SeqFold score		•									
PMF score		0.65	0.74	0.51	99.0	92.0	0.55	0.88	0.92	0.86	0.07
Verify score		0.1	0	-0.12	0.15	0.31	0.09	0.22	0.16	0.38	-0.81
PSI- BLAST		1.40E-32	3.40E-34	7.00E-07	1.70E-53	6.80E-54	3.40E-36	1.70E-60	3.40E-41	0.0065	0.00039
End		349	346	351	400	398	349	. 397	397	145	171
Start AA		147	146	258	146	141	147	147	143	17	146
Chain ID						٧			V	∢	A
PDB ID		lian	1jnk	1jnk	lkoa	1kob	1p38	1phk	1tki	1587	lega
SEQ NO:		965	596		965	965	296	596	596	601	602

PDB annotation	ENDOCYTOSIS/EXOCYTOSIS SYNAPTOTAGMIN ASSOCIATED 35 KDA PROTEIN, P35A, THREE HELIX BUNDLE	PHOSPHOTRANSFERASE RHOGAP DOMAIN; PHOSPHOTRANSFERASE, TPASE ACTIVATING PROTEIN, GAP, CDC42, 2 PHOSPHOINOSITIDE 3- KINASE, SH3 DOMAIN, SH2 DOMAIN, 3 SIGNAL TRANSDUCTION	PHOSPHOTRANSFERASE RHOGAP DOMAIN; PHOSPHOTRANSFERASE, TPASE ACTIVATING PROTEIN, GAP, CDC42, 2 PHOSPHONOSITIDE 3- KINASE, SH3 DOMAIN, SH2 DOMAIN, 3 SIGNAL TRANSDUCTION	PHOSPHOTRANSFERASE RHOGAP DOMAIN; PHOSPHOTRANSFERASE, TPASE ACTIVATING PROTEIN, GAP, CDCC42, 2 PHOSPHOINOSITIDE 3- KINASE, SH3 DOMAIN, SH2 DOMAIN, 3 SIGNAL TRANSDUCTION	G-PROTEIN CDC42 GTPASE- ACTIVATING PROTEIN; G-PROTEIN, GAP, SIGNAL-TRANSDUCTION	G-PROTEIN CDC42 GTPASE- ACTIVATING PROTEIN; G-PROTEIN, GAP, SIGNAL-TRANSDUCTION	COMPLEX(GTPASE ACTIVATIN/PROTO-ONCOGENE) GTPASE-ACTIVATING PROTEIN RHOGAP; COMPLEX (GTPASE ACTIVATION/PROTO-ONCOGENE), GTPASE, 2 TRANSITION STATE, GAP	COMPLEX(GTPASE ACTIVATN/PROTO-ONCOGENE) GTPASE-ACTIVATING PROTEIN RHOGAP; COMPLEX (GTPASE - ACTIVATION/PROTO-ONCOGENE), GTPASE, 2 TRANSITION STATE, GAP	COMPLEX(GTPASE
Coumpound	SYNTAXIN-1A; CHAIN: A, B, C;	PHOSPHATIDYLINOSITOL 3- KINASE; CHAIN: A, B;	PHOSPHATIDYLNOSITOL 3- KINASE; CHAIN: A, B;	PHOSPHATIDYLINOSITOL 3- KINASE; CHAIN: A, B;	RHOGAP; CHAIN: NULL;	RHOGAP; CHAIN: NULL;	P50-RHOGAP; CHAIN: A; TRANSFORMING PROTEIN RHOA; CHAIN: B;	P50-RHOGAP; CHAIN: A; TRANSFORMING PROTEIN RHOA; CHAIN: B;	P50-RHOGAP; CHAIN: A;
SeqFold score	-		·		112.2		118.67		
PMF	0.07	_	-	-		-		_	1
Verify score	-0.14	0.43	0.78	0.77		0.58		0.54	6.0
PSI- BLAST	0.0039	1.30E-20	. 1.30E-20	2.60E-40	1.80E-30	1.80E-30	9.10E-48	1.10E-30	9.10E-48
End	669	192	192	217	212	189	212	201	212
Start	260	25	25	25	13	16	15	16	16
Chain TO	¥	∢	g.	В			¥	∢	A
PDB TD	lez3	upp w	1pb · w	Ipb w	lrgp	lrgp	1tx4	1tx4	1¤4
SEQ NO.	909	605	905	\$09	605	909	605	509	605

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PDB annotation	ACTIVATN/PROTO-ONCOGENE) GTPASE-ACTIVATING PROTEIN RHOGAP; COMPLEX (GTPASE ACTIVATION/PROTO-ONCOGENE), GTPASE, 2 TRANSITION STATE, GAP	MERCURY DETOXIFICATION MERCURIC TRANSPORT PROTEIN; MERCURY DETOXIFICATION, PENPLASMIC, HEAVY METAL TRANSPORT, 2 ALPHA-BETA SANDWICH	HYDROLASE COPPER- TRANSPORTING ATPASE, COPPER- BINDING DOMAIN, HYDROLASE	NUCLEAR IMPORT RECEPTOR KARYOPHERIN ALPHA; NUCLEAR IMPORT RECEPTOR, NUCLEAR LOCALIZATION SIGNAL, 2 ARMADILLO REPEATS, AUTOINHIBITION, INTRASTERIC REGULATION	ARMADILLO REPEAT ARMADILLO REPEAT, BETA-CATENIN, CYTOSKELETON	COMPLEX (TRANSFERASE/PEPTIDE) COMPLEX (TRANSFERASE/PEPTIDE)	COMPLEX (TRANSFERASE/PEPTIDE) COMPLEX (TRANSFERASE/PEPTIDE)	COMPLEX (TRANSFERASE/PEPTIDE) ITAM PEPTIDE; COMPLEX (TRANSFERASE/PEPTIDE), SYK, KINASE, SH2 DOMAIN, ITAM
Coumpound	TRANSFORMING PROTEIN RHOA; CHAIN: B;	MERP; CHAIN: NULL;	MENKES COPPER- TRANSPORTING ATPASE; CHAIN: NULL;	IMPORTIN ALPHA; CHAIN: A;	BETA-CATENIN; CHAIN: NULL;	C-SRC TYROSINE KINASE; CHAIN: A, B; ACE-FORMYL PHOSPHOTYR-GLU-(N,N- DIPENTYL AMINE); CHAIN: C. D:	C-SRC TYROSINE KINASE; CHAIN: A, B: ACE-FORMYL PHOSPHOTYR-GLU-(N.N- DIPENTYL AMINE); CHAIN: C. D:	SYK KINASE; CHAIN: A, C, E, G, I, K; T-CELL SURFACE GLYCOPROTEIN CD3 EPSILON CHAIN; CHAIN: B, D, F, H, J, L;
SeqFold score						172.37		
PMF		pura.	_	0.46	0.96			-
Verify score		0.86	0.81	-0.02	0.42		1.07	0.57
PSI- BLAST		0.00026	0.0012	1.20E-06	0.00026	6.50E-41	6.50E-41	7.80E-33
End AA		203	203	91	148	255	255	255
Start AA		143	143	21	16	150	150	126
Chain ID				∢		4	٧	∢
PDB UD		lafi	law0	lial	3bct	1a09	1a09	1881
SEQ S B S		909	909	909	909	607	607	209

PDB annotation	COMPLEX (TRANSFERASE/PEPTIDE) ITAM PEPTIDE; COMPLEX (TRANSFERASE/PEPTIDE), SYK, KINASE, SH2 DOMAIN, ITAM			PROTEIN KINASE CDK2; PROTEIN KINASS, CELL CYCLE, PHOSPHORYLATION, STAUROSPORINE, 2 CELL DIVISION, MITOSIS, INHIBITION	COMPLEX (ISOMERASE/PROTEIN KINASE) FKBP12; SERINE/THREONING-PROTEIN
Coumpound	SYK KINASE; CHAIN: A, C, E, G, I, K; T-CELL SURFACE GLYCOPROTEIN CD3 EPSILON CHAIN; CHAIN: B, D, F, H, I, L;	TRANSFERASE(PHOSPHOTR ANSFERASE) \$C-IAMP\$-DEPENDENT PROTEIN KINASE (E.C.2.7.1.37) (\$C/APK\$) 1APM 3 (CATALYTIC SUBUNIT) ALPHA ISOENZYME MUTANT WITH SER 139 1APM 4 REPLACED BY ALA (\$1394\$) COMPLEX WITH THE PEPTIDE 1APM 5 INHIBITOR PKI(5-24) AND THE DETERGENT MEGA-8 1APM 6	TRANSFERASE(PHOSPHOTR ANSFERASE) SC-/AMP\$- DEPENDENT PROTEIN KINASE (E.C.2.7.1.37) (\$C/APLYTIC SUBUNIT) ALPHA ISOENZYME MUTANT WITH SER 139 1APM 4 REPLACED BY ALA (\$139A\$) COMPLEX WITH THE PEPTIDE 1APM \$ INHEBITOR PKI(\$-24) AND THE DETERGENT MEGA-8 1APM 6	CYCLIN-DEPENDENT PROTEIN KINASE 2; CHAIN: NULL;	FK506-BINDING PROTEIN; CHAIN: A, C, E, G; TGF-B SUPERFAMILY RECEPTOR
SeqFold score		97.15			102.06
PMF	6:0		-	_	
Verify score	0.26		0.81	0.34	
PSI- BLAST	1.30E-46	1.308-35	1.30E-35	1.30E-33	2.60E-74
End	372	542	\$15	529	533
Start AA	151	255	282	281	241
Chain	∢	м	ω		В
PDB	1881	qa u	qal e	laql	1b6c
SEQ SEQ	607	, 200	607	607	607

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PDB annotation	KINASE RECEPTOR R4; COMPLEX (ISOMERASE/PROTEIN KINASE), RECEPTOR 2 SERINE/THREONINE KINASE	COMPLEX (ISOMERASEPROTEIN KINASE) FKBP12; SERINETHREONINE-PROTEIN KINASE RECEPTOR R4; COMPLEX (ISOMERASEPROTEIN KINASE), RECEPTOR 2 SERINETHREONINE KINASE	V-SRC SH2 DOMAIN SRC SH2; V-SRC SH2 DOMAIN, PHOSPHOTYROSINE RECOGNITION DOMAIN, PP60 2 SRC SH2 DOMAIN	V-SRC SH2 DOMAIN SRC SH2; V-SRC SH2 DOMAIN, PHOSPHOTYROSINE RECOGNITION DOMAIN, PP60 2 SRC SH2 DOMAIN	COMPLEX (INHIBITOR PROTEIN/KINASE) INHIBITOR PROTEIN, CYCLIN-DEPENDENT KINASE, CELL CYCLE 2 CONTROL, ALPHABETA, COMPLEX (INHIBITOR PROTEIN/KINASE)	COMPLEX (INHIBITOR PROTEIN/KINASE) INHIBITOR PROTEIN, CYCLIN-DEPENDENT KINASE, CELL CYCLE 2 CONTROL, ALPHABETA, COMPLEX (INHIBITOR PROTEIN/KINASE)		
Coumpound	TYPE I; CHAIN: B, D, F, H;	FK506-BINDING PROTEIN; CHAIN: A, C, E, G, TGF-B SUPERFAMILY RECEPTOR TYPE I; CHAIN: B, D, F, H;	PP60 V-SRC TYROSINE KINASE TRANSFORMING PROTEIN; CHAIN: NULL;	PP60 V-SRC TYROSINE KINASE TRANSFORMING PROTEIN; CHAIN: NULL;	CYCLIN-DEPENDENT KINASE 6; CHAIN: A; P19INK4D; CHAIN: B;	CYCLIN-DEPENDENT KINASE 6; CHAIN: A; P19INK4D; CHAIN: B;	PHOSPHOTRANSFERASE CAMP-DEPENDENT PROTEIN KINASE CATALYTIC SUBUNIT ICMK 3 (E.C.2.7.1.37) ICMK 4	TRANSFERASE(PHOSPHOTR ANSFERASE) CAMP- DEPENDENT PROTEIN KINASE (E.C.2.7.1.37) (CAPK) ICTP 3 (CATALYTIC
SeqFold score				165.48		99.54		99.49
PMF		н					-	
Verify score		0.51	1.13		0.66		0.7	
PSI- BLAST		2.60B-74	2.60E-40	2.60E-40	1.205-34	1.20E-34	9.10E-35	6.50E-35
End		531	259	263	529	538	515	542
Start		256	155	155	269	270	282	. 722
Chain		æ	_		4	∢	ធ	មា
PDB		1b6c	16KI	15KI	1bix	1blx	lcm k	1сф
S B S		607	209	209	607	209	607	607

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PDB annotation				TRANSFERASE KINASE DOMAIN, AUTOINHIBITORY FRAGMENT, HOMODIMER	PROTEIN KINASE CDK2; TRANSFERASE, SERINETHREONINE PROTEIN KINASE, ATP-BINDING, 2 CELL CYCLE, CELL DIVISION, MITOSIS, PHOSPHORYLATION	PROTEIN KINASE CDK2; TRANSFERASE, SERINE/THREONINE PROTEIN KINASE, ATP-BINDING, 2 CELL CYCLE, CELL DIVISION, MITOSIS, PHOSPHORYLATION	TRANSFERASE INK3; TRANSFERASE, INK3 MAP KINASE, SERINE/THREONINE PROTEIN 2 KINASE	COMPLEX (TRANSFERASE/PEPTIDE) SRC, SH3 DOMAIN, LIGANDS, NON- PEPTIDE ELEMENTS, 2 COMPLEX (TRANSFERASE/PEPTIDE)	TRANSFERASE MITOGEN ACTIVATED PROTEIN KINASE; TRANSFERASE, MAP KINASE, SERNETHREOMNE-PROTEIN KINASE, 2 P38	
Coumpound	SUBUNIT) 1CTP 4	TRANSFERASE(PHOSPHOTR ANSFERASE) CAMP- DEPENDENT PROTEIN	KINASE (E.C.2.7.1.37) (CAPK) 1CTP 3 (CATALYTIC STRINGT) (CTP 4	SERINETHREONINE- PROTEIN KINASE PAK- ALPHA; CHAIN: A, B; SERINETHREONINE- PROTEIN KINASE PAK-	ALPHA; CHAIN; C; D; HUMAN CYCLN. DEPENDENT KINASE 2; CHAIN: NULL;	HUMAN CYCLIN- DEPENDENT KINASE 2; CHAIN: NULL;	C-JUN N-TERMINAL KINASE; CHAIN: NULL;	C-SRC; CHAIN: C; NL1 (MN7- MN2-MN1-PLPPLP); CHAIN: N;	MAP KINASE P38; CHAIN: NULL;	PHOSPHOTRANSFERASE V- SRC TYROSINE KINASE TRANSFORMING PROTEIN (PHOSPHOTYROSINE 1SHA 3 RECOGNITION DOMAIN
SeqFold score				·	111.67					165.79
PMF		-				-	i-		~	
Verify score		0.55		0.71		0.43	9.0	0.13	0.44	
PSI- BLAST		6.50E-35		1.30E-42	1.30E-37	1.30E-37	1.30E-33	2.60E-19	2.60E-32	9.10E-40
End		515		519	538	529	538	149	517	256
Start		282		272	273	281	269	88	269	154
Chain D		ம		U				ပ		4
PDB DD	1	Jcfp		1f3m	Ibcl	1hcl	ljnk	1nlo	1p38	Isha
SEQ EQ		209		209	607	607	607	607	607	209

PDB annotation		,	CYTOSKELETON CAPPING PROTEIN, CALCIUM-BINDING, DUPLICATION, REPEAT, 2 SH3 DOMAIN, CYTOSKELETON	CYTOSKELETON CAPPING PROTEIN, CALCIUM-BINDING, DUPLICATION, REPEAT, 2 SH3 DOMAIN, CYTOSKELETON	TRANSFERASE TRANSFERASE, TYROSINE KINASE, SH3, SH2, ONCOPROTEIN	TRANSFERASE TRANSFERASE, TYROSINE KINASE, SH3, SH2, ONCOPROTEIN	TYROSINE PHOSPHATASE SYP, SHPTP-2: TYROSINE PHOSPHATASE, INSULIN SIGNALING, SH2 PROTEIN	-	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC
Coumpound	SH2) (E.C.2.7.I.112) COMPLEX WITH 1SHA 4 PHOSPHOPEPTIDE A (TYR-VAL-PRO-MET-LEU, PHOSPHORYLATED TYR) 1SHA 5	PHOSPHOTRANSFERASE V- SRC TYROSINE KINASE TRANSFORMING PROTEIN 3 (PHOSPHOTYROSINE 1SHA 3 RECOGNITION DOMAIN SH2) (E.C.2.7.1.112) COMPLEX WITH 1SHA 4 PHOSPHOPEPTIDE A (TYR- VAL-PRO-MET-LEU, PHOSPHORYLATED TYR) 1SHA 5	ALPHA-SPECTRIN; CHAIN: NULL;	ALPHA-SPECTRIN; CHAIN: NULL;	ABL TYROSINE KINASE; CHAIN: NULL;	ABL TÝROSINE KINASE; CHAIN: NULL;	SHP-2; CHAIN: A, B;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX
SeqFold score					121.36				
PMF		-	0.89	0.01		-	0.54	0.31	0.74
Verify score		0.97	-0.02	-0.25		9.0	-0.13	-0.13	0
PSI- BLAST		9.10E-40	1.30E-14	3.90E-09	9.10E-34	9.10E-34	6.50E-59	9.10E-21	1.30E-32
End		256	166	130	255	255	425	284	290
Start AA	-	154	100	75	79	08	75	, 203	209
Chain		∢					4	< .	A
PDB ID		l sha	Ituc	Itud	2abi	2abl	2shp	1a1h	laih
SEQ SEQ		607	209	209	607	607	209	809	809

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PDB annotation	FINGER, DNA-BINDING PROTEIN								,				COMPLEX (ZINC FINGER/DNA) ZINC	FINGER, PROTEIN-DNA	CENTERACTION, PROTEIN DESIGN, 2	(ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC	FINGER, PROTEIN-DNA	INTERACTION, PROTEIN DESIGN, 2	CRYSTAL STRUCTURE, COMPLEX	COMPLEX (ZINC FINGER/DNA) ZINC	FINGER, PROTEIN-DNA	INTERACTION, PROTEIN DESIGN, 2	CRYSTAL STRUCTURE, COMPLEX	(ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC	FINGER, PROTEIN-UNA	CBYCTAI CTDICTIBE COMPLEX	(ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC	FINGER, PROTEIN-DNA INTER ACTION PROTEIN DESIGN 2	CRYSTAL STRUCTURE, COMPLEX
Coumpound	OI ICONITIC! EOTIDE	BINDING SITE; CHAIN: B, C;	TRANSCRIPTION REGULATION YEAST	TRANSCRIPTION FACTOR	ADR1 (RESIDUES 102 - 130)	1ARD 3 (AMINO TERMINAL	ANC THOER DOMESTIC	1ARD 4 (ADR1B) 1ARD 5	DNA-BINDING PROTEIN	BINDING PROTEIN MBP-1	MUTANT WITH CYS 11 1BBO	(C11ABU) (NMR, 60 STRIICTIMES) 18BO 4	DNA; CHAIN: A, B, D, E;	CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;		DNA; CHAIN: A, B, D, E;	CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;		DNIA: CHARM A B D F.	CONSENSIS ZINC FINGER	PROTEIN; CHAIN: C, F, G;			DNA; CHAIN: A, B, D, E;	CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C. F. C.		DNA; CHAIN: A, B, D, E;	CONSENSUS ZINC FINGER	rkolein; chain: c, r, g;
SeqFold										,					_																	
PMF	31036		0.05						0.17				0.87				_				-	-				-				_		
Verify	arnic.		-0.34						-0.3			-	0.16				0.32				9	6.07				19.0				0.62		
PSI-	Dimon		6.50E-05						2.60E-16				1.30E-22				6.50E-33				. 200	1.205-42				5.20E-44				9.10E-47		
End	*		537						537				283				311				,,,,	252				367				395		
Start	AA		509						485				202				230				0,0	607				286				314		
Chain	a												O	,			O	,				د				ပ				U		
PDB	a		lard						ogq.				Ime	>			t	>				e ;	<u> </u>			Ime	^			Ime	۸	
SEQ	ΒÖ		809		_				809				809				809					sõ S				809				809		

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PDB annotation	(ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINDERDINA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA FINTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINCEKUNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA CHEACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL, STRUCTURE, COMPLEX (ZINC FINGER/DNA)
Coumpound		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAÎN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAÎN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;
SeqFold score		102.94			!				
PMF			-		-		0.22	-	
Verify			0.74	0.53	0.33	0.05	-0.08	0.2	0.29
PSI- BLAST		9.10E-47	7.80E-46	1.30E-45	1.30E-40	2.60E-34	3.90E-32	3.90E-42	3.90E-46
End		396	423	451	479	505	589	617	645
Start		314	342	370		426	454	536	564
Chain		U	U	O	v	O	v	o .	O
PDB	+	Jme y	y y	Jme y	1me y	1me y	y y	Ime y	y y
SEQ	Ö	809	809	809	809	809	809	809	809

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PDB annotation	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	ZINC FINGEK IRANSCKIF IION FACTOR SPI; ZINC FINGER, TRANSCRIPTION ACTIVATION, SPI	ZINC FINGER IRANSCRIFTION FACTOR SP1, ZINC FINGER, TRANSCRIPTION ACTIVATION, SP1	COMPLEX (TRANSCRIPTION REGULATIONDNA) COMPLEX (TRANSCRIPTION REGULATIONDNA), RNA POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN		COMPLEX (TRANSCRIPTION REGULATIONDNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATIONDNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN
Coumpound	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	SP1F3; CHAIN: NULL;	SP1F2; CHAIN: NULL;	TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	TRANSCRIPTION REGULATION TRANSCRIPTIONAL ELONGATION FACTOR SII (TFIIS, NUCLEIC-ACID 1TFI 3 BINDING DOMAIN) (NMR, 12 STRUCTURES) 1TFI 4	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;
SeqFold					109.61			
PMF	1	-	10.0	0.13		0.24	0.03	1
Verify	0.33	0.6	-0.29	-0.23		-0.22	-0.34	0.26
PSI- BLAST	2.60E-46	2.60E-45	7.80E-05	9.10E-05	2.60E-71	0.0091	7.80E-29	6.50E-43
End	673	702	537	537	505	520	311	340
Start	592	620	509	509	342	480	165	229
Chain	U	U			¥		U	U
PDB TD	y y	y y	lsp1	1sp2	1116	1 2	Iubd	lubd
	ÖN 809	809	809	809	809	809	809	809

PDB annotation	RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATIONIDNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATIONIDNA)	COMPLEX (TRANSCRIPTION REGULATIONDNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATIONDNA) YING-YANG 1; REANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATIONDNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1: TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YYI, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG I; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YYI, ZINC 2
Coumpound		YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YY I; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;
SeqFold score							
PMF			 -	_		0.34	0.99
Verify score		0.55	0.58	0.26	0.34	-0.28	-0.04
PSI- BLAST		7.80E-52	1.30E-53	5.20E-53	9.10E-51	2.60E-39	5.20E-45
End		368.	395	451	479	589	819
Start AA		263	284	341	374	424	508
Chain ID		U	U	O	ပ	U	ပ
PDB ID		lubd	lubd	lubd	lubd	lubd	1ubd
SEQ	Ž	809	809	809	809	809	809

PDB annotation	FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INTIATION, INTIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	TRANSCRIPTION REGULATION TRANSCRIPTION REGULATION, ADRI, ZINC FINGER, NMR			COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLJ; GLJ, ZINC FINGER, COMPLEX (DNA-BINDING PROTEIN/DNA)		1-1
Coumpound		YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	ADRI; CHAIN: NULL;	COMPLEX(TRANSCRIPTION REGULATION/DNA) TRAMTRACK PROTEIN (TWO ZINC-FINGER PEPTIDE) COMPLEXED WITH 2DRP 3 DNA 2DRP 4	COMPLEX(TRANSCRIPTION REGULATION/DNA) TRAMTRACK PROTEIN (TWO ZINC-FINGER PEPTIDE) COMPLEXED WITH 2DRP 3 DNA 2DRP 4	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII;
SeqFold score								,	
PMF			•••	0.41	90.0	0.23		_	_
Verify		-0.07	0.06	-0.15	0.11	0.28	1.0	0.45	0.32
PSI- BLAST		5.20E-56	1.30E-54	2.60E-18	1.30E-15	6.50E-18	9.10E-58	1.00E-65	1.30E-67
End		674	701	209	530	563	369	397	480
Start		264	290	455	479	505	230	259	314
Chain		U	U		¥	4	∀	4	A
PDB ID		Iubd	1ubd	2adr	2drp	2drp	2gli	2gli	2gli
SEQ SEQ	į	809	809	809	809	809	809	809	809

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PDB annotation	PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLJ; GLJ, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)		COMPLEX (GTPASE-ACTIVATING/GTP-BINDING) COMPLEX (GTPASE-ACTIVATING/GTP-BINDING), GTPASE	TRANSPORT PROTEIN TC4; GTPASE, NUCLEAR TRANSPORT, TRANSPORT PROTEIN	TRANSPORT PROTEIN TC4; GTPASE, NUCLEAR TRANSPORT, TRANSPORT PROTEIN	SIGNALING PROTEIN PROTEIN- PROTEIN COMPLEX, ANTIPARALLEL COILED-COIL	SIGNALING PROTEIN PROTEIN- PROTEIN COMPLEX, ANTIPARALLEL COILED-COIL	SIGNALING PROTEIN P21-RAC2; RHO GDI 2, RHO-GDI BETA, LY-GDI; BETA
Coumpound	CHAIN: A, DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GL11; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;		P50-RHOGAP; CHAIN: A, B, C; CDC42HS; CHAIN: D, B, F;	GTP-BINDING PROTEIN RAN; CHAIN: A, B;	GTP-BINDING PROTEIN RAN; CHAIN: A, B;	HIS-TAGGED TRANSFORMING PROTEIN RHOA(0-181); CHAIN: A; PKN: CHAIN: B;	HIS-TAGGED TRANSFORMING PROTEIN RHOA(0-181); CHAIN: A; PKN; CHAIN: B;	RAS-RELATED C3 BOTULINUM TOXIN
SeqFold					92.71			55.55		61.64		
PMF score		0.37	-	0.99			_		0.66			
Verify score		0	0.3	0.16			0.46		0.15		0.19	0.14
PSI- BLAST		1.30E-56	1.30E-64	2.60E-70	2.60E-70		2.60E-20	3.90E-24	3.90E-24	1.30E-20	1.30E-20	2.60E-20
End AA		619	675	701	703		165	192	181	187	170	172
Start AA		398	536	564	564		9		9	_	<u>ن</u>	9
Chain ID		4	¥	4	∢		Ω	æ	В	4	4	. ⋖
PDB		2gli	2gli	2gli	2gli		1am 4	1byu	1byu	lcxz	lexz	1ds6
SEQ EQ	Ž	809	809	809	809		609	609	609	609	609	609

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PDB annotation	SANDWHICH, PROTEIN-PROTEIN-COMPLEX, G-DOMAIN, 2 IMMUNOGLOBULIN FOLD, WALKER FOLD, GTP-BINDING PROTEIN	G PROTEIN G PROTEIN, RAS, ARF, ARF6, MEMBRANE TRAFFIC BECTEN TRANSPORT GDP-BINDING.	MEMBRANE TRAFFICKIN, NON- MYRISTOYLATED 1HUR 16 MOTEUR TRAFFICKIN, NON- MOTEUR TRAFFICKING.	MEMBRANE TRAFFICKIN, NON- MYRISTOYLATED IHUR 16	SMALL GTPASE KAKY OFFERIN BETA, P95 SMALL GTPASE, NUCLEAR TRANSPORT RECEPTOR	BETA, P9S SMALL GTPASE, NUCLEAR TRANSPORT RECEPTOR	GTP-BINDING GIP-BINDING, OIFASE, SMALL G-PROTEIN, RHO FAMILY, RAS SUPER 2 FAMILY	GTP-BINDING GIP-BINDING, OIFASE, SMALL G-PROTEIN, RHO FAMILY, RAS SUPER 2 FAMILY GRAND FOR THE STANDING FARE		COMPLEX (SMALL OF ASSENDED FOR PROTEIN) COMPLEX (SMALL GTPASENUCLEAR PROTEIN), SMALL GTPASE, 2 NUCLEAR TRANSPORT	COMPLEX(GIPASE ACTIVATN/PROTO-ONCOGENE) GTPASE-ACTIVATING PROTEIN RHOGAP, COMPLEX (GTPASE ACTIVATION/PROTO-ONCOGENE), GTPASE, 2 TRANSITION STATE, GAP	COMPLEX(GIPASE ACTIVATMPROTO-ONCOGENE) GTPASE-ACTIVATING PROTEIN
Coumpound	SUBSTRATE 2; CHAIN: A; RHO GDP-DISSOCIATION INHIBITOR 2; CHAIN: B;	ADP-RIBOSYLATION FACTOR 6; CHAIN: A;	HUMAN ADP- RIBOSYLATION FACTOR 1; IHUR 5 CHAIN: A, B; IHUR 7	HUMAN ADP- RIBOSYLATION FACTOR 1; IHUR 5 CHAIN: A, B; IHUR 7	RAN; CHAIN: A, C; IMPORTIN BETA SUBUNIT: CHAIN: B, D;	RAN; CHAIN: A, C; IMPORTIN BETA SUBUNIT; CHAIN: B, D;	RAC1; CHAIN: NULL;	RACI; CHAIN: NULL;	RAN; CHAIN: A, C; NUCLEAR PORE COMPLEX PROTEIN NUP358; CHAIN: B, D;	RAN; CHAIN: A, C; NUCLEAR PORE COMPLEX PROTEIN NUP358; CHAIN: B, D;	Pso-Rhogap; Chain: A; Transforming protein Rhoa; Chain: B;	P50-RHOGAP; CHAIN: A; TRANSFORMING PROTEIN RHOA; CHAIN: B;
SeqFold			55.9		71.05		52.65		77.78		54.35	
PMF		0.57		0.55		66.0				0.6		-
Verify score		0.12		-0.12		0.04		0.36		-0.02		0.37
PSI- BLAST		1.20E-22	7.80E-23	7.80E-23	3.90E-23	3.90E-23	3.90E-21	3.90E-21	9.10E-24	9.10E-24	2.60E-20	2.60E-20
End	1	891	172	168	174	173	189	171	187	180	185	165
Start		9		9	3	9	2	9	8	9	7	~
Chain		4	4	V V	A	4			O	O	В	B
PDB		1e0s /	1hur	Ihur	1ibr	libr	dm 1	Į _	Іпр	Imp	11x4	1¤4
SEQ		609	609	609	609	609	609	609	609	609	609	609

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PDB annotation	RHOGAP, COMPLEX (GTPASE ACTIVATION/PROTO-ONCOGENE), GTPASE, 2 TRANSITION STATE, GAP	COMPLEX (INHIBITOR PROTEIN/KINASE) INHIBITOR PROTEIN, CYCLIN-DEPENDENT KINASE, CELL CYCLE 2 CONTROL, ALPHA/BETA, COMPLEX (INHIBITOR PROTEIN/KINASE)	PROTEIN KINASE CDK2; TRANSFERASE, SERINE/THREONINE PROTEIN KINASE, ATP-BINDING, 2 CELL CYCLE, CELL DIVISION, MITOSIS, PHOSPHORYLATION	UBIQUITIN CONJUGATION UBC2; UBIQUITIN CONJUGATION, UBIQUITIN-CONJUGATING ENZYME	LIGASE UBIQUITIN, UBIQUITIN- CONJUGATING ENZYME, YEAST	UBIQUITIN CONJUGATION UBCI; UBIQUITIN CONJUGATION, LIGASE	MIISCI E PROTEIN CTNC: CARDIAC.	MUSCLE PROTEIN, REGULATORY, CALCIUM BINDING	HYDROLASE CALCINEURIN; HYDROLASE, PHOSPHATASE, IMMUNOSUPPRESSION	CALCIUM-BINDING PROTEIN CALCIUM-MYRISTOYL SWITCH, CALCUIM-BINDING PROTEIN	CALCIUM-REGULATED MUSCLE CONTRACTION MUSCLE CONTRACTION, CALCIUM-BINDING, TROPONIN, E-F. HAND, 2 OPEN CONFORMATION REGULATORY DOMAIN, CALCIUM-REGULATED 3 MISCI E CONTRACTION	CALCIUM-BINDING PROTEIN EF-
Coumpound		CYCLIN-DEPENDENT KINASE 6; CHAIN: A; PI9INK4D; CHAIN: B;	HUMAN CYCLIN- DEPENDENT KINASE 2; CHAIN: NULL;	UBIQUITIN-CONJUGATING ENZYME RAD6; CHAIN: A, B, C;	UBIQUITIN CONJUGATING ENZYME; CHAIN: A;	UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	TEOPONIN C. CHAIN: NIII I	INOFORMA C, CITAMA: NOBE,	SERINE/THREONINE PHOSPHATASE 2B; CHAIN: A, B;	RECOVERIN; CHAIN: NULL;	TROPONIN C; CHAIN: NULL;	TROPONIN C; 1TNX 4
SeqFold score		129.83	118.75	54.69	54.87	59.33	70.39	V0.38	78.45	67.12	68.38	67.26
PMF												
Verify score												
PSI- BLAST		2.60E-52	7.80E-50	1.00E-15	2.60E-17	2.60E-14	א טעם על	3.20E-06	7.80E-21	5.20E-20	9.10E-17	1.20E-16
End		691	687	337	335	333	071	80	476	491	465	463
Start AA		370	371	167	168	191	21.0	31./	320	311	320	320
Chain		∢		. 4	V				ω			
PDB ID		1blx	1hcl	·layz	1909	2aak		18,4	1aui	1 iku	ltcf	1thx
SEQ	Š	612	612	628	628	628	[631	631	631	631	631

PDB annotation	HAND ITNX 14	PHOSPHOTRANSFERASE PHOSPHOTRANSFERASE	LOF BLUMPT COMPANY	PHOSPHOTRANSFERANSE NUCLEOSIDE TRIPHOSPHATE, NUCLEOSIDE DIPHOSPHATE INUE 10	COMPLEX (GTPASE-	ACTIVATING/GTP-BINDING) COMPLEX (GTPASE-ACTIVATING/GTP-BINDING), GTPASE ACTIVATION	SIGNALING PROTEIN GTP-BINDING PROTEINS, PROTEIN-PROTEIN COMPLEX, EFFECTORS	SIGNALING PROTEIN GTP-BINDING PROTEIN RHOA, GTPASE RHOA; RHO	GDI I, RHO GTPASE, G-PROTEIN, SIGNALING PROTEIN	SIGNALING PROTEIN G PROTEIN, GTP HYDROLYSIS, KINETIC CRYSTALLOGRAPHY, 2 SIGNALING PROTEIN	SIGNALING PROTEIN PROTEIN- PROTEIN COMPLEX, ANTIPARALLEL COILED-COIL	SIGNALING PROTEIN P21-RAC2; RHO GDI 2, RHO-GDI BETA, LY-GDI; BETA SANDWHICH, PROTEIN-PROTEIN COMPLEX, G-DOMAIN, 2 IMMUNOGLOBULIN FOLD, WALKER FOLD, GTP-BINDING PROTEIN
Coumpound	CHAIN: NULL; 1TNX 5	NUCLEOSIDE DIPHOSPHATE TRANSFERASE; CHAIN: A, B, C;	PHOSPHOTRANSFERASE NUCLEOSIDE DIPHOSPHATE KINASE (E.C.2.7.4.6) INSQ 3	NUCLEOSIDE DIPHOSPHATE KINASE; INUE 4 CHAIN: A, B, C, D, E, F; INUE 5	DEC BUCCAD: CUAIN: A B	C; CDC42HS; CHAIN: D, E, F;	RAS-RELATED PROTEIN RAP-1A; CHAIN: A; PROTO- ONKOGENE SERINE/THREONINE PROTEIN KINASE CHAIN: B;	TRANSFORMING PROTEIN BHOA: C. RHO	GDP DISSOCIATION INHIBITOR ALPHA; CHAIN: E, F;	TRANSFORMING PROTEIN P21/H-RAS-1; CHAIN: A;	HIS-TAGGED TRANSFORMING PROTEIN RHOA(0-181); CHAIN: A; PKN: CHAIN: B;	RAS-RELATED C3 BOTULINUM TOXIN SUBSTRATE 2; CHAIN: A; RHO GDP-DISSOCIATION INHIBITOR 2; CHAIN: B;
SeqFold score		105.78	104.27	105.45					~			·
PMF					0.00	ς; 	0	0.65		0.36	0.93	0.54
Verify score						0.31	0.17	0.24		0.23	0.39	0.05
PSI- BLAST		3.60E-53	3.60E-50	9.00E-53	,,,	3.60E-56	5.40E-56	9.00E-62		1.80E-59	9.00E-62	1.40E-62
End		115	115	116		263	264	27.1		265	264	267
Start AA.		-	_	-		88	98	68		98	88	88
Chain		4	4	∢ .		Д	¥	A		4	4	V
PDB	1	1be4	lnsq	lnue		lam 4	lely	1000		1ctg	1cxz	1ds6
SEQ	ÖŽ	646	646	646		658	658	658		658	658	658

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PDB annotation	PROTEIN TRANSPORT GDP-BINDING, MEMBRANE TRAFFICKIN, NON- MYRISTOYLATED 1HUR 16	GTP-BINDING GTP-BINDING, GTPASE, SMALL G-PROTEIN, RHO FAMILY, RAS SUPER 2 FAMILY	COMPLEX(GTPASE ACTIVATN/PROTO-ONCOGENE) GTPASE-ACTIVATING PROTEIN RHOGAP, COMPLEX (GTPASE ACTIVATION/PROTO-ONCOGENE), GTPASE, 2 TRANSITION STATE, GAP	HYDROLASE CDC42/CDC42GAP: CDC42/CDC42GAP; TRANSITION STATE, G-PROTEIN, GAP, CDC42, ALF3., HYDROLASE	HYDROLASE G PROTEIN, VESICULAR TRAFFICKING, GTP HYDROLYSIS, RAB 2 PROTEIN, NEUROTRANSMITTER RELEASE, HYDROLASE		TRANSCRIPTION REGULATION PROTO-ONCOGENE, NUCLEAR BODIES (PODS), LEUKEMIA, 2 TRANSCRIPTION REGULATION		LIGASE CBL, UBCH7, ZAP-70, E2, UBIQUITIN, E3, PHOSPHORYLATION, 2 TYROSINE KINASE, UBIQUITINATION, PROTEIN DEGRADATION,	METAL BINDING PROTEIN RING FINGER PROTEIN MATI; RING FINGER (C3HC4)	HELICASE DNA REPAIR, DNA
Coumpound	HUMAN ADP- RIBOSYLATION FACTOR 1; 1HUR 5 CHAIN: A, B; 1HUR 7	RAC1; CHAIN: NULL;	PSO-RHOGAP, CHAIN: A; TRANSFORMING PROTEIN RHOA; CHAIN: B;	GTP BINDING PROTEIN (G25K), CHAIN: A; GTPASE ACTIVATING PROTEIN (RHG), CHAIN: B;	RAB3A; CHAIN: A;		TRANSCRIPTION FACTOR PML; CHAIN: NULL;	VIRUS EQUINE HERPES VIRUS-1 (C3HC4, OR RING DOMAIN) ICHC 3 (NMR, 1 STRUCTURE) ICHC 4	SIGNAL TRANSDUCTION PROTEIN CBL; CHAIN: A; ZAP-70 PEPTIDE; CHAIN: B; UBIQUITIN-CONJUGATING ENZYME E12-18 KDA UBCH7; CHAIN: C;	CDK-ACTIVATING KINASE ASSEMBLY FACTOR MATI; CHAIN: A;	PCRA; CHAIN: NULL;
SeqFold score	54.59										
PMF		0.45	0.82	0.4	0.47		0.25	86.0	0.9	0.09	0.21
Verify score		0.19	0.31	-0.01	0.11		-0.73	0.24	0.25	-0.44	-0.75
PSI- BLAST	7.20E-11	1.30E-63	1.80E-58	7.20E-60	1.10E-54		9.00E-06	9.00E-15	1.805-10	1.30E-05	0.002
End	266	267	263	270	265		59	60	99	72	218
Start AA	27	88	68	88	98		91		81	18	172
Chain ID	∢		a	4	∢				¥	Ą	
PDB TD	1 hur	1 mh	1124	2ngr	3rab		1bor	1chc	1fbv	1825	1pjr
S B S	658	859		658	658		629	629	629	659	199

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PDB annotation	REPLICATION, SOS RESPONSE, HELICASE, 2 ATP-BINDING, DNA-	HELICASE DNA REPAIR, DNA	REPLICATION, SOS RESPONSE, HEI ICASE, ATP-2 BINDING, DNA-	BINDING	COMPLEX (HELICASE/DNA) COMPLEX (HELICASE/DNA),	HELICASE, DNA UNWINDING	HYDROLASE/DNA ATP-DEPENDENT	HELICASE PCRA; HELICASE PCRA,	HYDROLASE, DNA, PRODUCT	COMPLEX				TRANSCRIPTION HELIX-BUNDLE		FIND IN THE PROPERTY OF THE PR	I KANSONI IION IION IION			•											
Coumpound		BCB A (SUBINITY: CHAIN: A:	PCRA (SUBUNIT); CHAIN: B;	PCRA (SUBUNIT); CHAIN: D;	ATP-DEPENDENT DNA	HELICASE KEP; CHAIN: A, B, DNA CHAIN: C;	HELICASE PCRA; CHAIN: A,	F; HELICASE PCRA; CHAIN:	B, G, DINA (3- I)(*TP*TP*TP*TP*T)-3');	CHAIN: C, D; DNA (5'-	D(*GP*C)-3'); CHAIN: H; DNA	(5'-D(*AP*CP*TP*GP*C)-3');		TO ANICCIDITION	ELONGATION FACTOR S-II;	CHAIN: A;	TRANSCRIPTION	CHAIN: A:	TRANSCRIPTION	REGULATION	TRANSCRIPTIONAL	ELONGATION FACTOR SIL	PERDING DOMAIN ONE. 12	STRUCTURES) 1TF1 4	TRANSCRIPTION	REGULATION	TRANSCRIPTIONAL	ELONGATION FACTOR SIL	(TFIIS, NUCLEIC-ACID LIFTS	STRUCTURES) 1TF1 4	TRANSCRIPTION
SeqFold score																									84.88						
PMF		,	0.28		69.0		0.88								0.24		0.18		-	-					-						-
Verify			-0.84		-0.67		-0.31								-0.43		0.45		34.0	C .											0.45
PSI- BLAST	,		0.0045		0.00011		9 00E-06								1.00E-18		3.60E-12		20 100 .	1.305-22					2 00E 24	3.300-24					3.90E-24
End			202		238	}	236	2			_				64		64		١	332		_			,;;	766					332
Start			172		172	•	133	7/1							-		4			283					5	783					283
Chain			В			٠	1								4		4	.													
PDB	1		lqhg			agn i		2pjr							1eo0		1000	3		146						#					146
SEQ			199			8		199							663		199	3		663						663	_				663

PDB annotation		RNA BINDING PROTEIN/RNA XLRBPA; PROTEIN-RNA COMPLEX, DOUBLE STRANDED RNA, PROTEIN- RNA 2 INTERACTIONS, RNA-BINING PROTEIN, RNA BINDING	RNA BINDING PROTEIN/RNA XLRBPA; PROTEIN-RNA COMPLEX, DOUBLE STRANDED RNA, PROTEIN- MAY 2 INTERACTIONS, RNA-BINING PROTEIN, RNA BINDING PROTEIN/RNA	RNA BINDING PROTEIN/RNA XLRBPA; PROTEIN-RNA COMPLEX, DOUBLE STRANDED RNA, PROTEIN- RNA 2 INTERACTIONS, RNA-BINING PROTEIN, RNA BINDING PROTEIN/RNA	RNA BINDING PROTEIN/RNA XI,RBPA; PROTEIN-RNA COMPLEX, DOUBLE STRANDED RNA, PROTEIN- NAA 2 INTERACTIONS, RNA-BINING PROTEIN, RNA BINDING PROTEIN/RNA	RNÁ BINDING PROTEIN/RNA XLRBPA; PROTEIN-RNA COMPLEX, DOUBLE STRANDED RNA, PROTEIN- RNA 2 INTERACTIONS, RNA-BINING PROTEIN, RNA BINDING	CELL CYCLERNA DSRBDIII; NMR STRUCTURE, PROTEINRNA, PROTEIN DSRBD. DROSOPHILA, RNA 2 HAIRPIN
Coumpound	TRANSCRIPTIONAL ELONGATION FACTOR SII (TFIIS, NUCLEIC-ACID 1TFI 3 BINDING DOMAIN) (NMR, 12 STRUCTURES) 1TFI 4	DOUBLE STRANDED RNA BINDING PROTEIN A; CHAIN: A, B; RNA (5'- R(*GP*CP*CP*CP*CP*CP*CP*CP*CP*CP*CP*CP*CP*CP	DOUBLE STRANDED RNA BINDING PROTEIN A; CHAIN: A, B, RNA (5- I (*GP*GP*GP*CP*GP*CP F *GP*CP*C)-3'); CHAIN: C, D, E, G:	BLE STRANDED RNA ING PROTEIN A; IN: A, B; RNA (5* P*GP*CP*GP*CP*CP*CP*CP*CP*CP*CP*CP*CP*CP*CP*CP*CP	SLE STRANDED RNA ING PROTEIN A; N: A, B, RNA (5*- P*GP*CP*GP*CP*CP*CP*C)-3); CHAIN: C, D,	BLE STRANDED RNA ING PROTEIN A; N: A, B, RNA (5*- P*GP*CP*GP*CP*CP*CP*C)-3'; CHAIN: C, D,	ERNAL EFFECT EIN (STAUFEN); N: A; STAUFEN SLE-STRANDED RNA ING DOMAIN; CHAIN:
SeqFold score							
PMF		0.99	0.42	0.99	0.39	0.7	0.31
Verify score		0.06	0.45	0.11	-0.16	0.06	-0.12
PSI- BLAST		1.60E-13	1.30E-14	2.60E-16	1.80E-09	1.60E-07	1.80E-14
End		452		574	558	452	454
Start AA		388	390	512	514	388	
Chain 10		₹ .	<	<	<	a	V
PDB CD		Zipı	7 Jdi2	ZiPI	ZiP1	7 J	Ickz
SEQ NO.		664	664	664	664	664	999

PDB annotation		CELL CYCLERNA DSRBDIII; NMR STRUCTURE, PROTEINRNA, PROTEIN DSRBD, DROSOPHILA, RNA 2 HAIRPIN	CELL CYCLERNA DSRBDIII; NMR STRUCTURE, PROTEINRNA, PROTEIN DSRBD, DROSOPHILA, RNA 2 HAIRPIN	CELL CYCLERNA DSRBDIII; NMR STRUCTURE, PROTEINRNA, PROTEIN DSRBD, DROSOPHILA, RNA 2 HAIRPIN	TRANSFERASE DSRNA-BINDING DOMAIN, NMR, PKR, SOLUTION STRUCTURE, PROTEIN 2 KINASE, TRANSFERASE	TRANSFERASE DSRNA-BINDING DOMAIN, NMR, PKR, SOLUTION STRUCTURE, PROTEIN 2 KINASE, TRANSFERASE	DOUBLE STRANDED RNA BINDING DOMAIN STAUFEN ISTU 13	DOUBLE STRANDED RNA BINDING DOMAIN STAUFEN ISTU 13	DOUBLE STRANDED RNA BINDING DOMAIN STAUFEN ISTU 13	RNA-BINDING PROTEIN/RNA TRA PRE-MRNA; SPLICING REGULATION, RNP DOMAIN, RNA COMPLEX	GENE REGULATION/RNA POLY(A) BINDING PROTEIN 1, PABP 1; RRM,
Coumpound	B;	MATERNAL EFFECT PROTEIN (STAUFEN); CHAIN: A; STAUFEN DOUBLE-STRANDED RNA BINDING DOMAIN; CHAIN: B:	MATERNAL EFFECT PROTEIN (STAUFEN); CHANI: A; STAUFEN DOUBLE-STRANDED RNA BINDING DOMAIN; CHAIN: B;	MATERNAL EFFECT PROTEIN (STAUFEN); CHAIN: A; STAUFEN DOUBLE-STRANDED RNA BINDING DOMAIN; CHAIN: B;	PROTEIN KINASE PKR; CHAIN: A;	PROTEIN KINASE PKR; CHAIN: A:	MATERNAL EFFECT PROTEIN STAUFEN; 1STU 4	MATERNAL EFFECT PROTEIN STAUFEN; 1STU 4	MATERNAL EFFECT PROTEIN STAUFEN; 1STU 4	SXI-LETHAL PROTEIN; CHAIN: A, B; RNA (5'- R(P*GP*UP*UP*GP*UP*UP*U P*UP*UP*UP*UP*U-CHAIN: P, Q;	POLYDENYLATE BINDING PROTEIN 1; CHAIN: A, B, C,
SeqFold score											
PMF		69:0	0.39	-	0.06	60.0	0.07	-	0.11	0.92	-
Verify score		90:0-	0.04	0.75	-0.17	-0.05	0.23	0.19	-0.24	0.48	1.05
PSI- BLAST		5.20E-15	5.40E-05	1.30E-19	1.10E-15	7.20E-08	3.60E-13	3.90E-18	0.0009	1.30E-10	9.10E-11
End		451	557	574	469	558	454	575	557	8	88
Start		384	509	511	375	518	388	512	514	-	01
Chain		A	⋖	4	<	4				<	В
PDB UD		lekz	1ekz	lekz	1qu6	1qu6	lstu	1sta	1stu	167£	1cvj
SEQ NO.		664	664	964	664	664	664	664	664	999	999

										
PDB annotation	PROTEIN-RNA COMPLEX, GENE REGULATION/RNA	GENE REGULATIONRNA POLY(A) BINDING PROTEIN 1, PABP 1; RRM, PROTEIN-RNA COMPLEX, GENE REGULATION/RNA	RNA BINDING PROTEIN KNA- BINDING DOMAIN	STRUCTURAL PROTEIN PROTEIN C.S., RNP, RBD, RRM, RNA BINDING DOMAIN, NUCLEOLUS		RNA-BINDING DOMAIN RNA- BINDING DOMAIN, ALTERNATIVE SPLICING	RNA-BINDING PROTEIN SPLICING, UZ SNRNP, RBD, RNA-BINDING PROTEIN	COMPLEX (RIBONUCLEOPROTEIN/DNA) HNRNP A1, UP1; COMPLEX (RIBONUCLEOPROTEIN/DNA), HETEROGENEOUS NUCLEAR 2 RIBONUCLEOPROTEIN A1	TOXIN BINDING PROTEIN TWO DOMAINS: BETA PROPELLER AND AT DATA RECTA FOILD.	TOXIN BINDING PROTEIN TWO DOMAINS: BETA PROPELLER AND
Coumpound	D, E, F, G, H; RNA (5'- R(*AP*AP*AP*AP*AP*AP*AP *AP*AP*AP*A)-3'); CHAIN: M, N, O, P, Q, R, S, T;	POLYDENYLATE BINDING PROTEIN I: CHAIN: A, B, C, D, E, F, G, H; RNA (5'- R(*AP*AP*AP*AP*AP*AP*AP*AP *AP*AP*AP*AP*3); CHAIN: M, N, O, P, Q, R, S, T;	HU ANTIGEN C; CHAIN: A;	NUCLEOLIN RBDI; CHAIN: A;	RNA-BINDING PROTEIN SEX-LETHAL PROTEIN (C- TERMINUS, OR SECOND RNA-BINDING DOMAIN 1SXL 3 (RBD-2), RESIDUES: 199 - 294 PLUS N-TERMINAL MET) 1SXL 4 (NMK, 17 STRUCTURES) 1SXL 5	SEX-LETHAL PROTEIN; CHAIN: NULL;	SPLICING FACTOR U2AF 65 KD SUBUNIT; CHAIN: A;	HETEROGENEOUS NUCLEAR RIBONUCLEOPROTEIN A1; CHAIN: A; 12-NUCLEOTIDE SINGLE-STRANDED TELOMETRIC DNA; CHAIN: B;	TOLB PROTEIN; CHAIN: A;	TOLB PROTEIN; CHAIN: A;
SeqFold score										
PMF		-	66.0	0.64	8.0		-	-	6.04	0.39
Verify		1.03	0.84	0.56	0.72	1.18	0.89	0.87	0.18	0.21
PSI- BLAST		7.80E-11	1.30E-11	2.60E-11	1.30E-11	2.60E-11	7.80E-11	6.50E-11	0.0013	3.90E-06
End		88	88	68	88	88	88	88	269	217
Start			6	∞	9	6	6	8	126	11
Chain		н	¥	<			∢	∢	<	<
PDB		1cvj	149a	167	1sxl	2sxl	2u2f	2up1	lorz	lorz
SEQ	Ö	999	999	599	999	999	999	599	999	999

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PDB annotation	ALPHA/BETA FOLD	TRANSCRIPTION INHIBITOR BETA- PROPELLER	TRANSCRIPTION INHIBITOR BETA- PROPELLER	COMPLEX (GTP- BINDING/TRANSDUCER) BETA1, TRANSDUCIN BETA SUBUNIT; GAMMA1, TRANSDUCIN GAMMA SUBUNIT; COMPLEX (GTP- BINDING/TRANSDUCER), G PROTEIN, HETERO/TRIMER 2 SIGNAL TRANSDUCTION	COMPLEX (GTP- BINDING/TRANSDUCER) BETA!, TRANSDUCIN BETA SUBUNIT; GAMMA I, TRANSDUCIN GAMMA SUBUNIT; COMPLEX (GTP- BINDING/TRANSDUCER), G PROTEIN, HETEROTRUMER 2 SIGNAL TRANSDUCTION	COMPLEX (GTP- BINDING/TRANSDUCER) BETA!, TRANSDUCIN BETA SUBUNIT; GAMMA!, TRANSDUCIN GAMMA SUBUNIT; COMPLEX (GTP- BINDING/TRANSDUCER), G PROTEIN, HETEROTRUMER 2 SIGNAL TRANSDUCTION	COMPLEX (GTP- BINDING/TRANSDUCER) BETA1, TRANSDUCIN BETA SUBUNIT; GAMMA1, TRANSDUCIN GAMMA SUBUNIT; COMPLEX (GTP- BINDING/TRANSDUCER), G PROTEIN, HETEROTRIMER 2 SIGNAL TRANSDUCTION		TRANSFERASE DINUCLEOTIDE-
Coumpound		TRANSCRIPTIONAL REPRESSOR TUP1; CHAIN: A, B, C;	TRANSCRIPTIONAL REPRESSOR TUP1; CHAIN: A, B, C;	GT-ALPHA/GI-ALPHA CHIMERA; CHAIN: A; GT- BETA; CHAIN: B; GT- GAMMA; CHAIN: G;	GT-ALPHA/GI-ALPHA CHIMERA; CHAIN: A; GT- BETA; CHAIN: B; GT- GAMMA: CHAIN: G;	GT-ALPHA/GI-ALPHA CHIMERA; CHAIN: A; GT- BETA; CHAIN: B; GT- GAMMA; CHAIN: G;	GT-ALPHA/GI-ALPHA CHIMERA; CHAIN: A; GT- BETA; CHAIN: B; GT- GAMMA; CHAIN: G;		NICOTINATE
SeqFold score						91.76			
PMF score		0.84	69.0		-0.02		0.8		-0.18
Verify score		0.2	0.24	0.59	0.27		0.56		0.52
PSI- BLAST		1.80E-59	3.60E-69	7.20E-73	1.10E-57	1.10E-57	5.40E-56		7.80E-14
End AA		315	373	271	420	372	370		194
Start		1	30	П	118	59	71		6
Chain		¥.	V .	В	В	·α	Д		Ą
PDB ID		lerj	lerj	lgot	lgot	1got	1got		1d0s
SEQ	2	999	999	999	999	999	999	L	699

		_																		_										
PDB annotation	BINDING MOTIF, PHOSPHORIBOSYL TRANSFERASE		TRANSCRIPTION HELIX-BUNDLE	TRANSCRIPTION HELIX-BUNDLE																			TRANSCRIPTION REGULATION PROTO CONCOCENE MILCI EAD	BODIES (PODS) LEUKEMIA 2	TRANSCRIPTION REGULATION				LIGASE CBL, UBCH7, ZAP-70, E2,	UBIQUITIN, E3, PHOSPHORYLATION,
Coumpound	MONONUCLEOTIDE:5,6- CHAIN: A;		TRANSCRIPTION ELONGATION FACTOR S-II; CHAIN: A;	TRANSCRIPTION ELONGATION FACTOR S-II;	CHAIN: A;	TRANSCRIPTION	TRANSCRIPTIONAL	ELONGATION FACTOR SII	(TFIIS, NUCLEIC-ACID 1TFI 3	STRUCTURES) 1TF1 4	TRANSCRIPTION	REGULATION	TRANSCRIPTIONAL	ELONGATION FACTOR SII	DESIGNACESIC-ACID LIFE S	STRUCTURES) 1TF1 4	TRANSCRIPTION	REGULATION	TRANSCRIPTIONAL	ELONGATION FACTOR SII	(TFIIS, NUCLEIC-ACID 1TFI 3	STRUCTURES) 1TFI 4	TRANSCRIPTION FACTOR	rive, cream. Note,		VIRUS EQUINE HERPES	VIRUS-1 (C3HC4, OR RING	DOMAIN) ICHC 3 (NMR, 1	SIGNAL TRANSDUCTION	PROTEIN CBL; CHAIN: A;
SeqFold											75.24																			
PMF			0.24	0.21		1											_						0.01			9.0			0.45	
Verify score			-0.43	-0.03		0.24		_									0.26						-0.65			-0.48			-0.62	
PSI- BLAST			1.00E-18	1.10E-06		1.80E-21					1.30E-21						1.30E-21					3	1.40E-06			1.80E-13			9.10E-11	
End			2	09		257					257						257						342			352			351	
Start			-	4		211					212						212						300			302			277	
Chain ID			٧_	Y																									¥	
PDB CD			1000	1eo0		146					146						146						1bor			1chc			1fbv	
SEQ No do			11/9	671		671					671						129						672			672			219	

	_							
PDB annotation	2 TYRÖSINE KINASE, UBIQUITINATION, PROTEIN DEGRADATION,	LIGASE CBL, UBCH7, ZAP-70, E2, UBIQUITIN, E3, PHOSPHORYLATION, 2 TYROSINE KINASE, UBIQUITINATION, PROTEIN DEGRADATION,	APOPTOSIS INHIBITOR OF APOPTOSIS (IAP), NMR STRUCTURE, BACULOVIRAL 2 IAP REPEAT (BIR), ZINC BINDING DOMAIN	DNA-BINDING PROTEIN V(D)) RECOMBINATION ACTIVATING PROTEIN 1; RAG1, V(D)) RECOMBINATION, ANTIBODY, MAD, RING FINGER, 2 ZINC BINUCLEAR CLUSTER, ZINC FINGER, DNA- BINDING PROTEIN	DNA-BINDING PROTEIN V(D)J RECOMBINATION ACTIVATING PROTEIN I; RAGI, V(D)J RECOMBINATION, ANTIBODY, MAD, RING FINGER, 2 ZNC BINUCLEAR CLUSTER, ZNC FINGER, DNA- BINDING PROTEIN	PEPTIDE SYNTHETASE GRSA; PEPTIDE SYNTHETASE, GRSA,	PEPTIDE SYNTHETASE GRSA; PEPTIDE SYNTHETASE, GRSA, ADENYLATE FORMING	OXIDOREDUCTASE OXIDOREDUCTASE, MONOOXYGENASE, PHOTOPROTEIN, LUMINESCENCE
Coumpound	ZAP-70 PEPTIDE; CHAIN: B; UBIQUITIN-CONJUGATING ENZYME E12-18 KDA UBCH7; CHAIN: C;	SIGNAL TRANSDUCTION PROTEIN CBL; CHAIN: A; ZAP-70 PEFTIDE; CHAIN: B; UBIQUITIN-CONJUGATING ENZYME E12-18 KDA UBCH7; CHAIN: C;	INHIBITOR OF APOPTOSIS PROTEIN (2MIHB/C-IAP-1); CHAIN: A;	RAGI; CHAIN: NULL;	RAGI; CHAIN: NULL;	GRAMICIDIN SYNTHETASE 1; CHAIN: A, B; BUENIYI AI ANNE: CUAIN.	C, D; GRAMICIDIN SYNTHETASE I; CHAIN: A, B; PHENYLALANINE; CHAIN:	LUCIFERASE; CHAIN: NULL;
SeqFold score			_			158.05		173.59
PMF		0.55	0.37	8.0	0.84		_	
Verify score		-0.4	-0.73	0.34	-0.11		0.6	
PSI- BLAST	•	3.60E-13	0.0027	2.60E-09	7.20E-09	0	0	3.60E-93
End		349	270	344	342	576	571	574
Start AA		301	227		302	24	39	22
Chain ED		<	⋖		,	4	<	
PDB ID		1fbv	1qbh	Irmd	Irmd	lam u	lam u	1]ci
S B S		672	672	672	672	673	673	673

							_		_		_		_,		
PDB annotation	OXIDOREDUCTASE OXIDOREDUCTASE, MONOOXYGENASE, PHOTOPROTEIN, LUMINESCENCE	TRANSCRIPTION REGULATION PROTO-ONCOGENE, NUCLEAR BODIES (PODS), LEUKEMIA, 2 TRANSCRIPTION REGULATION	TRANSCRIPTION REGULATION PROTO-ONCOGENE, NUCLEAR	BODIES (PODS), LEUKEMIA, 2 TRANSCRIPTION REGULATION	TRANSCRPTION REGULATION PROTO-ONCOGENE, NUCLEAR BODIES (PODS), LEUKEMIA, 2 TRANSCRIPTION REGULATION	TRANSCRIPTION REGULATION PROTO-ONCOGENE, NUCLEAR BODIES (PODS), LEUKEMIA, 2 TRANSCRIPTION REGULATION		CELL MOTILITY PROTEIN MSP; CYTOSKELETAL PROTEIN, SPERM, CELL MOTILITY PROTEIN	CELL MOTH ITY BROTEIN MSB.	CYTOSKELETAL PROTEIN, SPERM, CELL MOTILITY PROTEIN		ENDOCYTOSIS/EXOCYTOSIS SYNAPTOTAGMIN ASSOCIATED 35 KDA ROTEIN, P35A, THREE HELIX BUNDLE		SCAFFOLD PROTEIN SCAFFOLD PROTEIN, PP2A, PHOSPHORYLATION, HEAT REPEAT	SMALL GTPASE KARYOPHERIN BETA, P95 SMALL GTPASE, NUCLEAR
Coumpound	LUCIFERASE; CHAIN: NULL;	TRANSCRIPTION FACTOR PML; CHAIN: NULL;	TRANSCRIPTION FACTOR PMI: CHAIN: NULL:		TRANSCRIPTION FACTOR PML; CHAIN: NULL;	TRANSCRIPTION FACTOR PML; CHAIN: NULL;		MAJOR SPERM PROTEIN; CHAIN: A, B;	. Contract of the contract of	MAJOK SPEKM PKOLEIN; CHAIN: A. B;		SYNTAXIN-1A; CHAIN: A, B, C;		PROTEIN PHOSPHATASE PP2A; CHAIN: A, B;	RAN; CHAIN: A, C; IMPORTIN BETA SUBUNIT;
SeqFold score															
PMF score	-	0.01	0.01		0.01	0.01		0.07		0.07		-0.19		-0.02	90.0
Verify score	0.77	0.21	0.21		0.21	0.21		-0.05		-0.03		0.12		0.16	0.02
PSI- BLAST	3.60E-93	0.0031	0.00054		0.0031	0.00054		7.80E-06		7.80E-06		7.80E-09		1.10E-51	5.40E-07
End	571	761	192		192	192		125		125		286		377	119
Start AA	36	144	144		144	144		34		34		165		-	22
Chain								4		∢		∢		4	Ø
PDB ID	11ci	1bor	1bor		1bor	1bor		lmsp		1msp		lez3		1b3u	1ibr
SEQ	673	675	675		919	929		619		089		189		682	682

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PDB annotation	TRANSPORT RECEPTOR	NUCLEAR TRANSPORT PROTEIN COMPLEX HEAT REPEATS, NUCLEAR TRANSPORT PROTEIN COMPLEX	ARMADILLO REPEAT ARMADILLO REPEAT, BETA-CATENIN, CYTOSKELETON		POLYRIBONUCLEOTIDE TRANSFERASE POLYNUCLEOTIDE PHOSPHORYLASE, GUANOSINE PLOYRIBONUCLEOTIDE TRANSFERASE, ATT: GTP 2 DIPHOSPHOTRANSFERASE, RNA PROCESSING, RNA DEGRADATION	POLYRIBONUCLEOTIDE TRANSFERASE POLYNUCLEOTIDE PHOSPHORYLASE, GUANOSINE POLYRIBONUCLEOTIDE TRANSFERASE, ATP-GTP DIPHOSPHOTRANSFERASE, 2 RNA PROCESSING, RNA DEGRADATION		OXIDOREDUCTASE OXIDOREDUCTASE, OXYGENREDUCTASE, DIRON- CENTRE, 2 FLAVOPROTEINS, LACTAMASE-FOLD	HYDROLASE METALLO-BETA- LACTAMASE, ANTIBIOTIC RESISTANCE, BINUCLEAR 2 ZINC, HYDROLASE	HYDROLASE METALLO-BETA- LACTAMASE, ANTIBIOTIC RESISTANCE, BINUCLEAR 2 ZINC, HYDROLASE	HYDROLASE HYDROLASE, BETA- LACTAMASE, ANTIBIOTIC, METALLOENZYME		DNA-BINDING PROTEIN RPA, OB-
Coumpound	CHAIN: B, D;	KARYOPHERIN BETA2; CHAIN: B; RAN; CHAIN: C;	BETA-CATENIN; CHAIN: NULL;		GUANOSINE PENTAPHOSPHATE SYNTHETASE; CHAIN: A;	GUANOSINE PENTAPHOSPHATE SYNTHETASE; CHAIN: A;		RUBREDOXIN:OXYGEN OXIDOREDUCTASE; CHAIN: A, B	PENICILINASE; CHAIN: A;	PENICILLINASE; CHAIN: A;	METALLO BETA- LACTAMASE II; CHAIN: A, B;		REPLICATION PROTEIN A 32
SeqFold score													
PMF		0.07	0.09		-	_		90:0-	0.04	0.09	-0.05		0.23
Verify score		0.11	-0.18		0.55	0.62		0.03	0.29	-0.22	0.05		0.01
PSI- BLAST		3.60E-37	1.30E-07		7.20E-58	7.20E-58		3.60E-13	1.30E-18	0.00052	3.60E-10		3.60E-38
End		400	363		273	273		191	198	558	187		181
Start		2	32			-		89	34	498	45		28
Chain		В			<	∢		∢	∢	4	∢		V
PDB CD	1	1qbk	3pct		163h	1e3p		1e5d	Isml	Isml	2pc2		lquq
SEQ	ë	289	682		683	683		684	684	684	684		989

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PDB annotation	FOLD, SSDNA-BINDING, DNA- BINDING PROTEIN	OXIDOREDUCTASE THIOREDOXIN M, THIOREDOXIN CH2, CHLOROPLASTIC THIOREDOXIN	OXIDOREDUCTASE DIMER, THIOREDOXIN, X-RAY CRYSTALLOGRAPHY, OXIDOREDUCTASE	ELECTRON TRANSPORT ELECTRON TRANSPORT	TRYPAREDOXIN TRYX-I; TRYPAREDOXIN, CRITHDIA FASCICULATA, THIOREDOXIN, 2 TRYPANOSOME, ANOMALOUS DISPERSION, OXIDATIVE STRESS, 3 OXIDOREDUCTASE	TRYPAREDOXIN TRYX-I; TRYPAREDOXIN, CRITHUIA FASCICULATA, THIOREDOXIN, 2 TRYPANOSOME, ANOMALOUS DISPERSION, OXIDATIVE STRESS, 3 OXIDOREDUCTASE	PEROXIDASE 2-CYS PEROXIREDOXIN, CALPROMOTIN PEROXIDASE, PEROXIREDOXIN, SULPHINIC ACID, THIOREDOXIN	OXIDOREDUCTASE HEME-BINDING PROTEIN 23 KD, HBP23; THIOREDOXIN FOLD, OXIDOREDUCTASE	ELECTRON TRANSPORT ALPHAMBETA OPEN-TWISTED PROTEIN, THIOL- DISULFIDE	T7 DNA POLYMERASE, DNA REPLICATION, NUCLEOTIDYL 2 TRANSFERASE, SEQUENCING, THIOREDOXIN, PROCESSIVITY FACTOR, 3 COMPLEX
Coumpound	KD SUBUNIT; CHAIN: A, C; REPLICATION PROTEIN A 14 KD SUBUNIT; CHAIN: B, D;	CHLOROPLAST THIOREDOXIN M CH2; CHAIN: A;	THIOREDOXIN; CHAIN: NULL;	THIOREDOXIN M; CHAIN: A, B;	TRYPAREDOXIN-1; CHAIN: A;	TRYPAREDOXIN-1; CHAIN: A;	HUMAN THIOREDOXIN PEROXIDASE-B; CHAIN: A, B, C, D, E, F, G, H, I, I;	THIOREDOXIN PEROXIDASE 2; CHAIN: A, B;	THIOREDOXIN, CHAIN: A;	DNA POLYMERASE; CHAIN: A; THIOREDOXIN; CHAIN: B; DNA; CHAIN: P, T;
SeqFold score					·					
PMF		0.11	0.09	0.01	0.18	0.34	-0.14	0.09	0.1	0.03
Verify		-0.28	0.08	-0.31	-0.87	0.32	0.06	0.1	-0.66	-0.36
PSI- BLAST		3.60E-13	3.60E-07	3.60E-11	5.40E-07	1.30E-12	5.40E-40	1.60E-40	1.30E-06	1.60E-12
End		147	142	143	127	181	233	210	187	150
Start AA		62	09	09	0/	75	48	48	78	62
Chain		V		4	4	e e	4	¥	V V	Ø
PDB CD		1dby	lerv	1 1 66	1qk8	1qk8	14m v	1992	lqu w	1t7р
SEQ	ğ	069	069	069	069	069	069	069	069	069

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PDB annotation	(HYDROLASE/ELECTRON TRANSPORT/DNA)				LIGASE CBL, UBCH7, ZAP-70, EZ, UBIQUITIN, E3, PHOSPHORYLATION, 2 TYROSINE KINASE, UBIQUITINATION, PROTEIN DEGRADATION,	LIGASE CBL, UBCH7, ZAP-70, E2, UBIQUITIN, E3, PHOSPHORYLATION, 2 TYROSINE KINASE, UBIQUITINATION, PROTEIN DEGRADATION,	DNA-BINDING PROTEIN V(D)) RECOMBINATION ACTIVATING PROTEIN 1; RAG1, V(D)) RECOMBINATION, ANTIBODY, MAD, RING FINGER, 2 ZNC BINUCLEAR CLUSTER, ZINC FINGER, DNA- BINDING PROTEIN	DNA-BINDING PROTEIN V(D)) RECOMBINATION ACTIVATING PROTEIN I; RAGI, V(D)I RECOMBINATION, ANTIBODY, MAD, RING FINGER, 2 ZINC BINUCLEAR CLUSTER, ZINC FINGER, DNA- BINDING PROTEIN	BETA.		TRANSCRIPTION INHIBITOR BETA-
Coumpound		ELECTRON TRANSPORT THIOREDOXIN 2TRXA 2	LIKAMS	VIRUS EQUINE HERPES VIRUS-1 (C3HC4, OR RING DOMAIN) 1CHC 3 (NMR, 1 STRUCTURE) 1CHC 4	SIGNAL TRANSDUCTION PROTEIN CBL; CHAÎN: A; ZAP-70 PEPTIDE; CHAÎN: B; UBIQUITIN-CONTUGATING ENZYME E12-18 KDA	SIGNAL TRANSDUCTION PROTEIN CBL; CHAIN: A; ZAP-70 PEPTIDE; CHAIN: B; UBIQUITIN-CONJUGATING ENZYME E12-18 KDA I IRCHT. CHAIN: C.	RAGI; CHAIN: NULL;	RAG1; CHAIN: NULL;		TRANSCRIPTIONAL REPRESSOR TUP1; CHAIN: A, B. C:	TRANSCRIPTIONAL
SeqFold score											
PMF		0.04		0.48	0.21	0.21	0.94	0.49		0.11	0.29
Verify		-0.1		-0.33	-0.58	-0.74	-0.07	-0.24		99.0	0.52
PSI- BLAST		1.60E-12		7.20E-13	1.20E-10	1.80E-06	1.00E-13	5.40E-06		5.40E-42	9.00E-51
End		150		363	363	368	363	368	-	307	369
Start		62		318	298	319.	297	319		4	71
Chain		4			4	₹				∢	▼
PDB	-+	2trx		Ichc	1fbv	1fbv	1rmd	1.md	1	lerj	[-
SEQ	Ö	069		169	1691	691	169	691		693	693

					<u> </u>		,		
PDB annotation	PROPELLER	COMPLEX (GTP. BINDING/TRANSDUCER) BETA1, TRANSDUCIN BETA SUBUNIT; GAMMA1, TRANSDUCIN GAMMA SUBUNIT; COMPLEX (GTP. BINDING/TRANSDUCER), G PROTEIN, HETEROTRIMER 2 SIGNAL TRANSDUCTION	COMPLEX (GTP-BINDING/TRANSDUCER) BETA1, TRANSDUCIN BETA SUBUNIT; GAMMAI, TRANSDUCIN GAMMA SUBUNIT; COMPLEX (GTP-BINDING/TRANSDUCER), G PROTEIN, TRANSDUCTION	COMPLEX (GTP- BINDING/TRANSDUCER) BETA1, TRANSDUCIN BETA SUBUNIT; GAMMA1, TRANSDUCIN GAMMA SUBUNIT; COMPLEX (GTP- BINDING/TRANSDUCER), G PROTEIN, HETEROTRUMER 2 SIGNAL TRANSDUCTION	TOXIN BINDING PROTEIN TWO DOMAINS: BETA PROPELLER AND ALPHA/BETA FOLD	TRANSCRIPTION INHIBITOR BETA- PROPELLER	TRANSCRIPTION INHIBITOR BETA- PROPELLER	TRANSCRIPTION INHIBITOR BETA- PROPELLER	TRANSCRIPTION INHIBITOR BETA- PROPELLER
Coumpound	REPRESSOR TUP1; CHAIN: A, B, C;	GT-ALPHA/GI-ALPHA CHIMERA; CHAIN: A; GT- BETA: CHAIN: B; GT- GAMMA; CHAIN: G;	GT-ALPHA/GI-ALPHA CHIMERA; CHAIN: A; GT- BETA; CHAIN: B; GT- GAMMA; CHAIN: G;	GT-ALPHA/GI-ALPHA CHIMERA; CHAIN: A; GT- BETA; CHAIN: B; GT- GAMMA; CHAIN: G;	TOLB PROTEIN; CHAIN: A;	TRANSCRIPTIONAL REPRESSOR TUP1; CHAIN: A, B. C.	TRANSCRIPTIONAL REPRESSOR TUP1; CHAIN: A, B. C.	TRANSCRIPTIONAL REPRESSOR TUP1; CHAIN: A, B, C;	TRANSCRIPTIONAL REPRESSOR TUP1; CHAIN: A,
SeqFold		70.14	·						
PMF score			0.16	-0.17	0.29	0.94	-	0.99	0.16
Verify score			0.39	0.26	0.2	0.12	0.46	10.01	0.27
PSI- BLAST		3.60E-57	3.60E-57	1.40E-36	9.00E-08	5.40E-78	1.80E-65	9.00E-57	1.40E-52
End AA		376	370	304	377	416	313	442	224
Start AA		28	39	v .	133	108	<u>8</u>	195	5
Chain D		В	ø.	Ω.	4	4	<	«	Ą
PDB U		lgot	1got	1got	Icrz	lerj	1erj	lerj	lerj
SEQ B B S		693	693	693	694	694	694	694	694

PDB annotation		COMPLEX (GTP-BINDICER) BETA1, BINDING/TRANSDUCER) BETA1, TRANSDUCIN BETA SUBUNIT; GAMMA1, TRANSDUCIN GAMMA SUBUNIT; COMPLEX (GTP-BINDING/TRANSDUCER), G PROTEIN, HETEROTRIMER 2 SIGNAL TRANSDUCTION	COMPLEX (GTP-BINDICER) BETA1, BINDING/TRANSDUCER) BETA1, TRANSDUCIN BETA SUBUNIT; GAMMAA1, TRANSDUCIN GAMMA SUBUNIT; COMPLEX (GTP-BINDING/TRANSDUCER), G PROTEIN, HETEROTRIMER 2 SIGNAL TRANSDUCTION	COMPLEX (GTP- BINDING/TRANSDUCER) BETA1, TRANSDUCIN BETA SUBUNIT; GAMMA1, TRANSDUCIN GAMMA SUBUNIT; COMPLEX (GTP- BINDING/TRANSDUCER), G PROTEIN, HETEROTRIMER 2 SIGNAL TRANSDUCTION	COMPLEX (GTP- BINDING/TRANSDUCER) BETA!, TRANSDUCIN BETA SUBUNIT; GAMMA!, TRANSDUCIN GAMMA SUBUNIT; COMPLEX (GTP- BINDING/TRANSDUCER), G PROTEIN, HETEROTRIMER 2 SIGNAL TRANSDUCTION	COMPLEX (GTP- BINDINGTRANSDUCER) BETA1, TRANSDUCIN BETA SUBUNIT; GAMMA1, TRANSDUCIN GAMMA SUBUNIT; COMPLEX (GTP- BINDINGTRANSDUCER), G PROTEIN, HETEROTRIMER 2 SIGNAL TRANSDUCTION
Coumpound	B, C;	GT-ALPHA/GI-ALPHA CHIMERA; CHAIN: A; GT- BETA; CHAIN: B; GT- GAMMA; CHAIN: G;	GT-ALPHA/GI-ALPHA CHIMERA; CHAIN: A; GT- BETA; CHAIN: B; GT- GAMMA; CHAIN: G;	GT-ALPHA/GI-ALPHA CHIMERA; CHAIN: A; GT- BETA; CHAIN: B; GT- GAMMA; CHAIN: G;	GT-ALPHA/GI-ALPHA CHIMERA; CHAIN: A; GT- BETA; CHAIN: B; GT- GAMMA; CHAIN: G;	GT-ALPHA/GI-ALPHA CHIMERA; CHAIN: A; GT- BETA; CHAIN: B; GT- GAMMA; CHAIN: G;
SeqFold score			105.8			
PMF		-		0.21	0.22	-
Verify		0.54		0.26	0.36	0.69
PSI- BLAST		5.40E-64	7.20E-81	3.60E-50	1.80E-53	7.20E-81
End		393	353	443	267	353
Start AA		103	12	190	4	26
Chain		m	а	a	m	В
PDB ID		1got	lgot	1got	1got	1got
SEQ NO.		694	694	694	694	694

				_							_		_	_		_	Τ			Т	_			_			Τ	_		_				
PDB annotation	() () () () ()	COMPLEX (ZINC FINGER/DINA)	COMPLEX (ZINC FINGERSES); FINGER, DNA-BINDING PROTEIN	(A KOLOED (DNA)	COMPLEX (ZINC FINGEROUNA) 7INC	COMPLEX (ZINC FINGENDING), ZINC FINGER, DNA-BINDING PROTEIN	11110	COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGEROUNS), ZINC FINGER DNA-BINDING PROTEIN		COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA), ZINC	FINGER, DNA-BINDING I NO LEIN	CONTRACTOR PINGER/DNA)	COMPLEX (ZINC FINGER/DNA), ZINC	FINGER, DNA-BINDING PROTEIN	(NAC) and the second	COMPLEX (ZINC FINGENDINA)	COMPLEX (ZINC FINGERODIA), ZINC	FINGEN, DISTRIBUTION OF	COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA), ZINC	FINGER, DNA-BINDING PROTEIN	COMPLEX (7INC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA), ZINC	FINGER, DNA-BINDING PROTEIN							COMPLEX (ZINC FINGER/DNA) ZINC	
Coumpound '		OGSR ZINC FINGER	PEPTIDE; CHAIN: A; DUPLEX	BINDING SITE; CHAIN: B, C;	OGSR ZINC FINGER	PEPTIDE; CHAIN: A; DUPLEX	BINDING SITE: CHAIN: B, C;	OGSR ZINC FINGER	PEPTIDE; CHAIN: A; DUPLEX	OLIGONUCLEO INDE	BINDING STILL, STEELS ST	PEPTIDE, CHAIN: A; DUPLEX	OLIGONUCLEOTIDE	BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE: CHAIN: A: DUPLEX	OLIGONUCLEOTIDE	BINDING SITE; CHAIN: B, C;	OGSR ZINC FINGER	PEPTIDE; CHAIN: A; DUPLEX	OLIGONUCLEOTIDE PRIDING SITE: CHAIN: B. C.	BINDING STE, CIT. C. S. S.	PEPTIDE; CHAIN: A; DUPLEX	OLIGONUCLEOTIDE	BINDING SITE, CHAIN: B, C,	QGSR ZINC FINGER PEPTIDE: CHAIN: A: DUPLEX	OLIGONUCLEOTIDE	BINDING SITE; CHAIN: B, C;	DNA-BINDING PROTEIN	HUMAN ENHANCER-	MITTANT WITH CYS 11 1BBO	3 REPLACED BY ABU	(C11ABU) (NMR, 60	DNA: CHAIN: A B D E:	MA, 944
SeqFold		00 78	```																														-	
PMF						_		50	6.0			0.81			69.0			110	7.7			69.0			0.39			0.09						0.45
Verify score	1	+				91.0		十	 51.9-			0.03			0			0.13	CI :0			0.14			-0.1			-0.35					- 1	-0.09
PSI-			1.30E-31		1	3.60E-27 (7	1.30E-31			7.20E-30			1.80E-28			2 400	5.40E-10			5.40E-28			3.60E-26			6 50F-11	:					1.80E-36
End		\dashv	282		+	309		ᅥ	337		_	421			480			1	86			554			154			264	<u> </u>					168
Start			200			228			228			341			397				9			484			74				± 					101
Chain			4			V			4			V			■	:			Ą			4			V		_							ပ
PDB	3	1	lalh		_	lalh			lalh			lalh			1alh				lalh			1a1h			laih				1000				_	Ime
SEQ		+	269			169	_		269			169			203	6			697	•		169			169	_			697					697

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PDB annotation	FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC PINGER, DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX	(CANCE TINGENDINA) COMPLEX (CANCETTION) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CXYSTAL STRUCTURE, COMPLEX (77NC PINCEP DNA)	FORTER CANDERS OF THE STRUCK FORTER OF THE STRUCK FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX	CONCENTRATION OF THE STRUCTURE OF THE STRUCTURE OF THE STRUCTURE, PROTEIN DESIGN, 2 INTERACTION, PROTEIN DESIGN, 2 CYNYSTAL STRUCTURE, COMPLEX CONCENTRATION	COMPLEX CONTROL CONTROL CONTROL CONTROL CONTROL CONTROL CONTROL CONTROL CONTROL COMPLEX CONTROL COMPLEX CONTROL COMPLEX CONTROL CONTROL COMPLEX CONTROL CONTRO	COMPLEX CINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 COXYSTAL STRUCTURE, COMPLEX CONTRESTED COMPLEX	COMPLEX CINC ENGENDARY COMPLEX CINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 COMPLEX	(ZINC FINGERIDNA) COMPLEX (ZINC FINGERIDNA) ZINC FINGER PROTEIN, DNA
Coumpound	CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA: CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E. CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E: CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER
SeqFold score					97.32				
PMF		0.16	0.8	-		96.0	0.22	0.95	0.99
Verify score		-0.26	-0.15	0.31		0.27	-0.23	0.21	0.41
PSI- BLAST		1.30E-16	9.00E-35	5.40E-47	5.40E-47	3.60E-46	1.30E-33	3.60E-47	1.40E-49
End		224	252	280	281	309	337	365	393
Start AA		102	156	199		227	227	283	312
Chain D		ပ	υ	ပ	 U	၁	၁	O	၁
PDB ID	۲.	Ime y	Ime y	Ime y				у	l me
SEQ ID NO:		697	169		_				697

PDB annotation	INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2	(ZINC FINGERDNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA	INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC	INTERACTION, PROTEIN DESIGN, 2	CRYSTAL STRUCTURE, COMPLEX (ZINC FINGEXDNA)	COMPLEX (ZINC FINGER/DNA) ZINC	INTERACTION, PROTEIN DESIGN, 2	CRYSTAL STRUCTURE, COMPLEX	(ZINC FINGERUNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA	INTERACTION, PROTEIN DESIGN, 2	CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC	FINGER, PROTEIN-DNA	CRYSTAL STRUCTURE, COMPLEX	(ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA	INTERACTION, PROTEIN DESIGN, 2	CRYSTAL STRUCTURE, COMPLEX	COMPLEX (ZINC FINGER/DNA) ZINC	FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2
Coumpound	PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E;	PROTEIN; CHAIN: C, F, G;		DNA; CHAIN: A, B, D, E;	PROTEIN: CHAIN: C. F. G.			DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;		DNA; CHAIN: A, B, D, E,	CONSENSUS ZINC FINGER	rectelly, chain, c, r, c,		DNA; CHAIN: A, B, D, E;	PROTEIN, CHAIN: C, F, G;		DNA: CHAIN: A. B. D. E.	CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;
SeqFold score																			•					
PMF		0.95		0.88		0.04			0.62			,	0			98.0							0.98	
Verify score		0.09		0		-0.68			0.03				-0.13			0.22				0.31	_		0.18	
PSI- BLAST		1.30E-48		1.80E-48		1.30E-21			3.60E-47				1.60E-35			3.60E-48				3.60E-50			1.10E-44	
End		422		451		208			480				126.			808		_		536			557	
Start		340		368		368			396				42			425				455			483	}
Chain ID		၁		C		ပ			C				ပ		•	O				ပ			C)
PDB		y Jmc		lme v	`	e .	>		Ime	^	,		I Be	_		Ime	<u>></u>			lme ,			JE I	у
SEQ September		269		169		269			697				697			697				269			269	

PDB annotation	CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGENDNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (TRANSCRIPTION REGULATIONDNA) TFIIIA; 5S GENE; NMR, TFIIIA, PROTEIN, DNA, TRANSCRIPTION FACTOR, 5S RNA 2 GENE, DNA BINDING PROTEIN, ZINC FINGER, COMPLEX 3 (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATIONIDNA) COMPLEX (TRANSCRIPTION REGULATIONIDNA), RNA POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATIONDNA) COMPLEX (TRANSCRIPTION (TRANSCRIPTION POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION
Coumpound	•	DNA; CHAIN: A, B, D, B; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN: CHAIN: C, F, G;	TRANSCRIPTION FACTOR IIIA; CHAIN: A; 3S RNA GENE; CHAIN: E, F;	TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;
SeqFold score						107.54		
PMF		0.58	0.94	-0.13	0.4		0.92	0.75
Verify		-0.2	0.53	0.3	-0.1		-0.08	-0.02
PSI- BLAST		1.80E-43	5.40E-13	1.10E-09	I.10E-19	2.60E-50	3.60E-37	3.60E-34
End		154	536	86	417	366	346	461
Start		73	509	71	341	199	200	313
Chain		ပ	5	O	<	∢	v	∢
PDB		Jme y	y y	Jme y	5	1466	<u>#</u>	<u>1</u> 16
SEQ PO	į	269	.697.	169	697	169	697	697

PDB annotation	INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATIONDNA) COMPLEX (TRANSCRIPTION PEGIT ATTOMMAN NA BNA	POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX	(IRANSCRIPTION RNA REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION	INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX	(TRANSCRIPTION	POLYMERASE III, 2 TRANSCRIPTION	INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION	TRANSCRIPTION INTILATION.	INITIATOR ELEMENT, YY1, ZINC 2	FINGER PROTEIN, DNA-PROTEIN	RECOGNITION, 3 COMPLEX	(TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION	TRECOLATION/DIA) TING-TANG 1,	INITIATOR ELEMENT, YYI, ZINC 2	FINGER PROTEIN, DNA-PROTEIN	RECOGNITION, 3 COMPLEX	(TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGILI ATTON/DNA) VING-YANG 1:	TRANSCRIPTION INITIATION.	INITIATOR ELEMENT, YYI, ZINC 2	FINGER PROTEIN, DNA-PROTEIN	RECOGNITION, 3 COMPLEX	COMPLEX (TRANSCRIPTION
Coumpound		TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;		TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE;	CHAIN: B, C, E, F;		TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE;	CHAIN: B, C, E, F;			YY1; CHAIN: C; ADENO-	NITIATOR FLEMENT DNA:	CHAIN: A, B;				YY1; CHAIN: C; ADENO-	ASSOCIATED VIRUS PS	CHAIN: A. B.	` ` `			YYI; CHAIN: C; ADENO-	INITIATOR ELEMENT DNA:	CHAIN: A. B.			YY1; CHAIN: C; ADENO-
SeqFold score																												
PMF		0.23		0.72			0.01				69.0						0.51						0.95					0.28
Verify score		-0.13		-0.02			-0.39				-0.35						-0.29						-0.18					-0.36
PSI- BLAST		5.40E-36		1.80E-30			1.80E-27				1.80E-20						1.30E-39		-				1.30E-31					1.30E-32
End		489		558			233				224						338						337					393
Start		341		426			74				104						204						235					260
Chain		4		<			V				ပ						ر د						ပ					O
PDB		116		<u>1</u>			9,111,0				1 ubd			_			Inpq						lubd					lubd
SEQ EQ	5	697		697			269				697						697						697					697

PDB annotation	REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INTIATION, INITATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN FRECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATIONDNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATIONDNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATIONIDNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATIONDA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)
Coumpound	ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;	YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;
SeqFold score				•		83.84
PMF		0.7	66'0	-	0.51	
Verify score		-0.3	-0.03	-0.11	0	
PSI- BLAST		1.80E-33	9.00E-35	2.60E-28	3.60E-32	1.80E-34
End		365	422	450	208	537
Start		263	320	345	404	427
Chain D		ပ	U	U	o.	υ
PDB ID		lubd	lubd	lubd	Inpq	lubd
SEQ EQ	Ö	169	169	169	697	

	1; 2 NA)	l; 2 NA)	1; 2 NA)	1; 2 3NA)			GLI,
PDB annotation	COMPLEX (TRANSCRIPTION REGULATIONDNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN DNA-RROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG I; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YYI, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG I; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YYI, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	TRANSCRIPTION REGULATION TRANSCRIPTION REGULATION, ADRI, ZINC FINGER, NMR		COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
Соитроипд	YY I; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	ADRI; CHAIN: NULL;	COMPLEX(TRANSCRIPTION REGULATION/DNA) TRAMTRACK PROTEIN (TWO ZINC-FINGER PEPTIDE) COMPLEXED WITH 2DRP 3 DNA 2DRP 4	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;
SeqFold score			·				
PMF score	0.88	0.46	0.35	0.07	0.31	-0.12	0.45
Verify score	0.06	-0.05	-0.25	-0.45	-0.15	0.26	-0.08
PSI- BLAST	1.80E-34	3.60E-32	1.30E-26	1.30E-18	2.60E-14	1.30E-15	7.20E-26
End	536		154	253	260	564	279
Start	433	463	69	78	512	208	101
Chain	U	U	ပ	ပ		¥	Ą
PDB DD	lubd	lubd	lubd	1ubd	2adr	2drp	2gli
SEQ B B B	697	169	697	697	269	697	269

											Н
Coumpound	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLI1; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII;
SeqFold score		ì			89.2						
PMF score	0.99	-	0.25	66.0		0.78	0.86	0.92	0.22	0.03	0.13
Verify score	-0.17	0.14	-0.24	0.06		-0.02	0.04	0.2	0.02	-0.05	-0.45
PSI- BLAST	6.50E-40	7.20E-28	5.20E-48	1.80E-32	5.20E-48	5.40E-35	3.60E-34	5.40E-33	1.60E-31	1.10E-24	3.90E-30
End	282	308	394	367	397	422	449	535	558	153	253
Start	102	198	200	235	255	292	320	404	.433	99	77
Chain	«	₹	¥	4	V	4	4	«	4	∢	A
PDB	2gli	. 2gli	2gli	2gli	2gli	2gli	2gli	2gli	2gli	2gli	2gli
SEQ NO DE	692	697	269	269	697	697	697	697	697	697	697

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PDB annotation	PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	THE PROPERTY OF THE PARTY OF TH	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1	AND 2, ECAD 12; CADHENIN, CELE ADHESION PROTEIN, CALCIUM PROTFIN	CELL ADHESION PROTEIN	EPITHELIAL CADHERIN DOMAINS 1	AND 2, ECAD12; CADHERIN, CELL	ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL	CELL A DIFFERION PROTEIN CELL	ADHESION PROTEIN	CELL ADHESION UVOMORULIN;	ADHESION	CELL ADHESION UVOMORULIN;	CADHERIN, CALCIUM BINDING, CELL ADHESION		TRANSFERASE TRANSFERASE, KINASE, CALVIN CYCLE	TRANSFERASE TRANSFERASE, KINASE, CALVIN CYCLE	TRANSFERASE UPRTASE;	TRANSFERASE, GLYCOSYLTRANSFERASE, UPRTASE	TRANSFERASE UPRTASE;	TRANSFERASE, GI YOOSYI TRANSFERASE, UPRITASE	TRANSFERASE UPRTASE;	TRANSFERASE,	TRANSFERASE PANK; PROTEIN-
Coumpound	CHAIN: A; DNA; CHAIN: C, D;	G	E-CADHERIN; CHAIN: A, B;		E-CADHERIN: CHAIN: A. B.				N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	A CANIMON! CITABLA.	N-CADHEKIN; CHAIN: A;	EPITHELIAL CADHERIN;	CHAIN: NULL;	EPITHELIAL CADHERIN;	CHAIN: NULL;		PHOSPHORIBULOKINASE; CHAIN: NULL;	PHOSPHORIBULOKINASE; CHAIN: NULL:	URACIL	PHOSPHORIBOSYLTRANSFE RASE CHAIN: D.C. B. A:	URACIL	PHOSPHORIBOSYLTRANSFE	IRACII.	PHOSPHORIBOSYLTRANSFE	PANTOTHENATE KINASE;
SeqFold score					7 69	r:				64.18									6.99	191.89						
PMF			0.63						-0.12			9.16	0.04		10.0			9.0				_		-		96.0
Verify score			-0.1						40.0			-0.03	-0.22		-0.08			0.17		L		0.84		2,0	3	0.43
PSI- BLAST			7.20E-32		7 200 22	7.2UE-32			1.40E-34	1.80E-32		1.80E-32	1.30E-06		1.30E-08			1.30E-13	1.30E-13	2.60E-73		3 60F-44		2 KOE-73		1.80E-35
End			302		200	2005			178	302		302	62		182			281	366	532		532		533	<u> </u>	294
Start AA			101		1	*			9	70		4	23		74	:		128	25	306		322		327	5	75
Chain			Ą			∢			V	¥		٧								ρ		2	1	6	a	_
PDB ID			ledh			legn			Incj	Incj		1ncj	Isuh		İsuh			la7j	1a7j	1643		1,543		16.43	3	lesm
SEQ	Ö		869			869			869	869		869	869		869	}		701	101	701		Ş		Ş	<u> </u>	5

PDB annotation	INHIBITOR COMPLEX	TRANSFERASE PANK; PROTEIN- INHIBITOR COMPLEX		PHOSPHOTRANSFERASE ADK; PHOSPHOTRANSFERASE, ZINC FINGER	KINASE KINASE, PHOSPHOTRANSFERASE	CHAPERONE HSP40; CHAPERONE, HEAT SHOCK, PROTEIN FOLDING, DNAK	CHAPERONE HSP40; CHAPERONE, HEAT SHOCK, PROTEIN FOLDING, DNAK	GENE REGULATION/RNA POLY(A) BINDING PROTEIN 1, PABP 1; RRM, PROTEIN-RNA COMPLEX, GENE REGULATION/RNA	GENE REGULATION/RNA POLY(A) BINDING PROTEIN 1, PABP 1; RRM, PROTEIN-RNA COMPLEX, GENE REGULATION/RNA	GENE REGULATION/RNA POLY(A) BINDING PROTEIN 1, PABP 1; RRM, PROTEIN-RNA COMPLEX, GENE REGULATION/RNA	RNA BINDING PROTEIN RNA- BINDING DOMAIN	ENDOCYTOSIS/EXOCYTOSIS SYNAPTOTAGMIN ASSOCIATED 35
Coumpound	CHAIN: A, B, C, D;	PANTOTHENATE KINASE; CHAIN: A, B, C, D;	TRANSFERASE URIDYLATE KINASE (E.C.2.7.4) COMPLEXED WITH ADP AND AMP IUKZ 3	ADENYLATE KINASE; CHAIN: NULL;	THYMIDYLATE KINASE; CHAIN: A, B, C, D, E, F, G, H;	DNAJ; CHAIN: NULL;	DNAJ; CHAIN: NULL;	POLYDENYLATE BINDING PROTEIN I; CHAIN: A, B, C, D, E, F, G, H; RNA (5'- R(*AP*AP*AP*AP*AP*AP*AP* *AP*AP*AP*A); CHAIN: M, N, O, P, O, R, S, T;	POLYDENYLATE BINDING PROTEIN I; CHAIN: A, B, C, D, E, F, G, H; RNA (5'- R(*AP*AP*AP*AP*AP*AP*AP*AP *AP*AP*AP*A); CHAIN: M, N, O, P, Q, R, S, T;	POLYDENYLATE BINDING PROTEIN I; CHAIN: A, B, C, D, B, F, G, H; RNA (5'- R(*AP*AP*AP*AP*AP*AP*AP*AP*AP *AP*AP*AP*A); CHAIN: M, N, O, P, O, R, S, T;	HU ANTIGEN C; CHAIN: A;	SYNTAXIN-1A; CHAIN: A, B, C;
SeqFold score			_			54.61						
PMF		66'0	0.54	0.21	0.03		0.52	0.69	0.46	0.46	0.93	-0.13
Verify score		0.55	0.19	-0.33	-0.38		0.52	0.07	0.13	0.03	0.15	0.14
PSI- BLAST		6.50E-67	1.20E-15	0.0078	1.30E-23	1.30E-31	1.30E-31	3.60E-14	1.80E-12	1.80E-12	1.80E-12	1.30E-09
End		303	302	302	303	84	84	255	259	259	251	186
Start AA		16	94	105	96	∞	6	182	182	182	185	77
Chain ID		¥			ပ			∢	ĹĽ,	Ħ	Y	A
PDB ID		lesm	lukz	1zin	3tmk	1bq0	1bq0	lcvj	lcvj	lcvj	148z	lez3
SEQ NO.		701	701	701	701	702	702	702	702	702	702	702

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PDB annotation	KDA PROTEIN, P35A, THREE HELIX BUNDLE	NUCLEAR PROTEIN HETEROGENEOUS NUCLEAR RIBOUNCLEOPROTEIN A1, NUCLEAR PROTEIN, HINNIP, RBD, RRM, RNP, RNA BINDING, 2 RIBONUCLEOPROTEIN	RNA BINDING PROTEIN RNA- BINDING DOMAIN	MOLECULAR CHAPERONE HDJ-1;	MOLECULAR CHAPERONE HDI-1; MOLECULAR CHAPERONE	MOLECULAR CHAPERONE HDJ-1; MOLECULAR CHAPERONE	RNA BINDING PROTEIN RNA- BINDING DOMAIN	COMPLEX (RIBONUCLEOPROTEIN/DNA) HNRNP A1, UP1; COMPLEX (RIBONUCLEOPROTEIN/DNA), HETEROGENEOUS NUCLEAR 2 RIBONUCLEOPROTEIN A1	COMPLEX (RIBONUCLEOPROTEIN/DNA) HNRNP A1, UP1; COMPLEX (RIBONUCLEOPROTEIN/DNA), HETEROGENEOUS NUCLEAR 2 RIBONUCLEOPROTEIN A1	ENDOCYTOSIS/EXOCYTOSIS SYNAPTOTAGMIN ASSOCIATED 35 KDA PROTEIN, P35A, THREE HELIX BUNDLE	
Coumpound		HNRNP A1; CHAIN: NULL;	HETEROGENEOUS NUCLEAR RIBONUCLEOPROTEIN DO;	HUMAN HSP40; CHAIN: NULL:	HUMAN HSP40; CHAIN: NULL;	HUMAN HSP40; CHAIN: NULL;	MUSASHII; CHAIN: A;	HETEROGENEOUS NUCLEAR REGONOCLEOPROTEIN A1; CHAIN: 4, 12-NUCLEOTIDE SINGLE-STRANDED TELOMETRIC DNA; CHAIN: B;	HETEROGENEOUS NUCLEAR RIBONUCLEOPROTEIN A1; CHAIN: A: 12-NUCLEOTIDE SINGLE-STRANDED TELOMETRIC DNA; CHAIN: B;	SYNTAXIN-1A; CHAIN: A, B, C;	
SeqFold score					56.45				·	·	
PMF score		0.19	0.51	1		0.47	0.18	0	0.05	0.11	
Verify score		0.33	0.12	0.57		0.01	0.42	0.16	-0.11	0.15	
PSI- BLAST		1.305-22	5.40E-16	2.60E-19	1.80E-28	1.80E-28	7.20E-17	5.40E-26	1.30E-18	6.50E-06	1,000
End AA		249	249	78	84	84 ·	249	255	259	575	Ş
Start AA		182	182	=	∞	8	182	165	182	468	
Chain D			4				A .	¥ .	∢	4	-
PDB TD		Ihal	Ihdi	1hdj	1hdj	[þ4]	2mss	2up1	2up1	lez3	
SEQ NO:		702	702	702	702	702	702	702	702	705	202

PDB annotation	REGULATION/DNA) GABPALPHA; GABPBETAI; COMPLEX (TRANSCRPTION TEQULATION/DNA), DNA-BINDING, 2 NUCLEAR PROTEIN, ETS DOMAIN, ANK YRIN REPEATS, TRANSCRIPTION 3 FACTOR	COMPLEX (TRANSCRIPTION REGULATION/DIA) GABPALPHA; GABPBETA1; COMPLEX (TRANSCRIPTION REGULATION/DIA), DNA-BINDING, 2 NUCLEAR PROTEIN, ETS DOMAIN, ANKYRIN REPEATS, TRANSCRIPTION 3 FACTOR	COMPLEX (TRANSCRIPTION REGULATION/DNA) GABPALPHA; GABPBETA1; COMPLEX (TRANSCRIPTION REGULATION/DNA), DNA-BINDING, 2 NUCLEAR PROTEIN, ETS DOMAIN, ANKYRIN REPEATS, TRANSCRIPTION 3 FACTOR		TUMOR SUPPRESSOR TUMOR SUPPRESSOR, CDK4/6 INHIBITOR, ANKYRIN MOTIF	TUMOR SUPPRESSOR TUMOR SUPPRESSOR, CDK4/6 INHIBITOR, ANKYRIN MOTIF	TUMOR SUPPRESSOR TUMOR SUPPRESSOR, CDK4/6 INHIBITOR, ANKYRIN MOTIF	TUMOR SUPPRESSOR TUMOR
Coumpound	ALPHA; CHAIN: A; GA BINDING PROTEIN BETA I; CHAIN: B; DNA; CHAIN: D, E;	GA BINDING PROTEIN ALPHA; CHAIN: A; GA BINDING PROTEIN BETA 1; CHAIN: B; DNA; CHAIN: D, E;	GA BINDING PROTEIN ALPHA; CHAIN: A; GA BINDING PROTEIN BETA I; CHAIN: B; DNA; CHAIN: D, E;	GA BINDING PROTEIN ALPHA; CHAIN: A; GA BINDING PROTEIN BETA I; CHAIN: B; DNA; CHAIN: D, E;	P19INK4D CDK4/6 INHIBITOR; CHAIN: NULL;	P19INK4D CDK4/6 INHIBITOR; CHAIN: NULL;	P19INK4D CDK4/6 INHIBITOR; CHAIN: NULL;	P19INK4D CDK4/6
SeqFold score			·		62.42			
PMF			-	0.86			_	-0.14
Verify score		0.84	0.49	0.51		0.85	0.72	0.03
PSI- BLAST		1.30E-30	1.80E-36	7.20E-33	3.60E-31	3.60E-31	9.00E-31	3.60E-27
End		273	323	351	298	293	320	224
Start		169	175	509	44	145	178	57
Chain D		æ	m	m				
PDB ID		lawc	lawc	lawc	1bd8	1pd8	1bd8	1 bd8
SEQ D	Ž	706	706	706	706	706	706	706

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PDB annotation	SUPPRESSOR, CDK4/6 INHIBITOR, ANKYRIN MOTIF	COMPLEX (INHIBITOR PROTEIN/KINASB) INHIBITOR PROTEIN, CYCLIN-DEPENDENT KINASE, CELL CYCLE 2 CONTROL, ALPHABETA, COMPLEX (INHIBITOR BOCTENIKNASE)	COMPLEX (INHIBITOR PROTEIN/KINASE) INHIBITOR PROTEIN, CYCLIN-DEPENDENT RINASE, CELL CYCLE 2 CONTROL, ALPHABETA, COMPLEX (INHIBITOR PROTEIN/KINASE)	COMPLEX (INHIBITOR PROTEIN/KINASE) INHIBITOR PROTEIN, CYCLIN-DEPENDENT KINASE, CELL CYCLE 2 CONTROL, ALPHABETA, COMPLEX (INHIBITOR PROTEIN/KINASE)	COMPLEX (INHIBITOR PROTEIN/KINASE) INHIBITOR PROTEIN, CYCLIN-DEPENDENT KINASE, CELL CYCLE 2 CONTROL, ALPHABETA, COMPLEX (INHIBITOR PROTEIN/KINASE)	HORMONE/GROWTH FACTOR P18- INK4C; CELL CYCLE INHIBITOR, P18INK4C; TUMOR, SUPPRESSOR, CYCLIN-2 DEPENDENT KINASE, HORMONE/GROWTH FACTOR	HORMONE/GROWTH FACTOR P18- DNK4C; CELL CYCLE INHIBITOR, P18INK4C, TUMOR, SUPPRESSOR, CYCLIN-2 DEPENDENT KINASE, HORMONE/GROWTH FACTOR	SIGNALING PROTEIN HELIX-TURN- HELIX, ANKYRIN REPEAT	CELL CYCLE INHIBITOR P18- INK4C(INK6); CELL CYCLE
Coumpound	INHIBITOR; CHAIN: NULL;	CYCLIN-DEPENDENT KINASE 6; CHAIN: A; P19INK4D; CHAIN: B;	CYCL'N-DEPENDENT KINASE 6; CHAIN: A; P19INK4D; CHAIN: B;	CYCLIN-DEPENDENT KINASE 6; CHAIN: A; P19INK4D; CHAIN: B;	CYCLIN-DEPENDENT KINASE 6; CHAIN: A; P19INK4D; CHAIN: B;	CYCLIN-DEPENDENT KINASE 6 INHIBITOR; CHAIN: A;	CYCLIN-DEPENDENT KINASE 6 INHIBITOR; CHAIN: A;	CYCLIN-DEPENDENT KINASE 4 INHIBITOR B; CHAIN: A;	CYCLIN-DEPENDENT KINASE 6 INHIBITOR;
SeqFold score		63.41				74.38			68.42
PMF			_	96.0	-0.13		0.95	_	
Verify score			0.71	0.65	0.24		0.61	0.55	
PSI- BLAST		1.10E-30	1.80E-30	1.10E-30	3.60E-26	9.00E-34	9.00E-34	1.30E-28	5.40E-33
End		297	293	315	224	302	329	273	296
Start AA		144	145	178		134	175	171	138
Chain 'UD		æ	æ	В	а	4	· 4	₹	Ą
PDB		1bix	1blx	1blx	1blx	1bu9	15u9	149s	1 ihb
SEQ SEQ	2	706	706	902	706	706	706	706	706

PDB annotation	REG/ANK REPEAT) COMPLEX (TRANSCRIPTION REGULATION/ANK REPEAT), ANK YRIN 2 REPEAT HELIX	COMPLEX (ANTI- ONCOGENEANKYRIN REPEATS) P53BP2; ANKYRIN REPEATS, SH3, P53, TUMOR SUPPRESSOR, MULTIGENE 2 FAMILY, NUCLEAR PROTEIN, PHOSPHORYLATION, DISEASE MUTATION, 3 POLYMORPHISM, COMPLEX (ANTI- ONCOGENE/ANKYRIN REPEATS)	CALCIUM-BINDING PROTEIN CALSEQUESTRIN, CALCIUM-BINDING PROTEIN, SARCOPLASMIC 2 RETICULUM, RABBIT SKELETAL MUSCLE	CALCIUM-BINDING PROTEIN CALSEQUESTRIN, CALCIUM-BINDING PROTEIN, SARCOPLASMIC 2 RETICULUM, RABBIT SKELETAL MUSCLE	ELECTRON TRANSPORT ELECTRON	TRANSPORT, REDOX-ACTIVE CENTER, ISOMERASE, 2 ENDOPLASMIC RETICULUM	OXIDOREDUCTASE THIOREDOXIN M, THIOREDOXIN CH2, CHLOROPLASTIC THIOREDOXIN	OXIDOREDUCTASE THIOREDOXIN M, THIOREDOXIN CH2, CHLOROPLASTIC THIOREDOXIN	OXIDOREDUCTASE DIMER, THIOREDOXIN, X-RAY CRYSTALLOGRAPHY, OXIDOREDUCTASE	ELECTRON TRANSPORT ELECTRON TRANSPORT	ELECTRON TRANSPORT ELECTRON TRANSPORT, REDOX-ACTIVE
Coumpound	C; NF-KAPPA-B P50; CHAIN: B, D; I-KAPPA-B-ALPHA; CHAIN: E, F;	P53; CHAIN: A; 53BP2; CHAIN: B;	CALSEQUESTRIN; CHAIN: NULL	CALSEQUESTRIN; CHAIN: NULL	PROTEIN DISULFIDE	ISOMERASE; CHAIN: NULL;	CHLOROPLAST THIOREDOXIN M CH2; CHAIN: A;	CHLOROPLAST THIOREDOXIN M CH2; CHAIN: A;	THIOREDOXIN; CHAIN: NULL;	THIOREDOXIN M; CHAIN: A, B;	PROTEIN DISULFIDE ISOMERASE; CHAIN: NULL;
SeqFold score		75.09	87.98				52.47				82.03
PMF				96.0	0.07			0.95	0.98	_	
Verify score				0.15	-0.03			0.3	0.54	0.81	
PSI- BLAST		1.80E-25	5.40E-42	5.40E-42	1 60E-13		5.40E-24	5.40E-24	1.10 E-23	9.00E-25	1.10E-31
End	,	362	336	332	230	ì	131	131	128	130	134
Start		170	61	20	133	}	23	30	14	27	50
Chain 10		œ					V	∢		∢	
PDB ID		lycs	1a8y	la8y	1 Piv	5	ldby	1dby	lerv	1fb6	lme k
SEQ EQ	Ž	706	709	709	96	}	709	709	709	709	402

PDB annotation	CENTER, ISOMERASE, 2 ENDOPLASMIC RETICULUM	ELECTRON TRANSPORT ELECTRON TRANSPORT, REDOX-ACTIVE CENTER, ISOMERASE, 2 ENDOPLASMIC RETICULUM	ELECTRON TRANSPORT ALPHA/BETA OPEN-TWISTED PROTEIN, THIOL- DISULFIDE	T7 DNA POLYMERASE, DNA REPLICATION, NUCLEOTIDYL 2 TRANSFERASE, SEQUENCING, THIOREDOXIN, PROCESSIVITY FACTOR, 3 COMPLEX (FYDROLASELECTRON TRANSPORTIDNA)	T7 DNA POLYMERASE, DNA REPLICATION, NUCLEOTIDYL 2 TRANSFERASE, SEQUENCING, THIOREDOXIN, PROCESSIVITY FACTOR, 3 COMPLEX (HYDROLASFELECTRON TRANSPORT/DNA)	ELECTRON TRANSPORT THIOREDOXIN 2; 1THX 1 OXIDO- REDUCTASE 1THX 16	ELECTRON TRANSPORT THIOREDOXIN 2; 1THX 7 OXIDO- REDUCTASE 1THX 16	ELECTRON TRANSPORT HTRX, HCH1, CH1; OXIDOREDUCTASE, ELECTRON TRANSPORT			DNA INTEGRATION
Coumpound		PROTEIN DISULFIDE ISOMERASE; CHAIN: NULL;	THIOREDOXIN; CHAIN: A;	DNA POLYMERASE; CHAIN: A; THIOREDOXIN; CHAIN: B; DNA; CHAIN: P, T;	DNA POLYMERASE; CHAIN: A; THIOREDOXIN; CHAIN: B; DNA; CHAIN: P, T;	THIOREDOXIN; ITHX 5 CHAIN: NULL; ITHX 6	THIOREDOXIN; ITHX 5 CHAIN: NULL; ITHX 6	THIOREDOXIN H; CHAIN: NULL;	ELECTRON TRANSPORT THIOREDOXIN 2TRXA 2 2TRXA 3	ELECTRON TRANSPORT THIOREDOXIN 2TRXA 2 2TRXA 3	AVIAN SARCOMA VIRUS INTEGRASE; 1ASU 7 CHAIN:
SeqFold score				60.72		55.88			62.2		
PMF		0.98	_		_		1	99.0		_	0.39
Verify score		0.43	0.62		0.39		0.75	0.33		0.39	-0.13
PSI- BLAST		1.10E-31	3.60E-24	1.60E-24	1.60E-24	9.10E-22	9.10E-22	9.00E-23	5.40E-25	5.40E-25	1.80E-25
End		133	130	128	130	. 131	125	129	131	130	283
Start AA		21	30	26	27	24	32	25	24	25	130
Chain 1D			A	æ	a				∢	4	
PDB		k Inc	19u w	177p	1t7p	1thx	1thx	1tof	2trx	2trx	lasu
SEQ	Š	709	709	709	709	709	709	709	709	709	715

								_		_	_	<u>.</u>			_					_			_	т-			_			_	_		_	\neg
PDB annotation	Comment of the second	TRANSFERASE DNA INIEGRATION	TRASFERASE DNA INTEGRATION, TRASFERASE	DNA INTEGRATION DNA	INTEGRATION, AIDS, POLYPROLEIN,	HYDROLASE, 2 ENDONOCLEASE,	POLYNOCEGOID IE INCHASTERATE, DNA RINDING 3 (VIRAL)	THE ANCEDD ACT INTEGRASE ROLLS	SARCOMA VIRUS, HIV. X-RAY	CRYSTALLOGRAPHY, 2 PROTEIN	STRUCTURE, TRANSFERASE	VIRUS/VIRAL PROTEIN INTEGRASE,	ROUS SARCOMA VIRUS, HIV, A-RAI	PROTEIN	TRANSFERASE MIXED BETA-SHEET	SURROUNDED BY ALPHA-HELICES	VIRUS/VIRAL PROTEIN HIV-1	INTEGRASE, POLITNOCEEOTIDIE	PROTEIN, DD35E	HYDROLASE DNA INTEGRATION,	INTEGRASE, HIV, HYDROLASE,	ASPARTYL 2 PROTEASE,	ENDONOCLEASE	- TANE CELL CVCI E ANTI-	ONCOGENE, REPEAT, ANK REPEAT	ANTI-ONCOGENE CELL CYCLE, ANTI-	ONCOGENE, REPEAT, ANK REPEAT	COMPLEX (TRANSCRIPTION	RECOLATION/DIAM) CABI ALL IM,	(TRANSCRIPTION	REGULATION/DNA), DNA-BINDING, 2	NUCLEAR PROTEIN, ETS DOMAIN,	3 FACTOR	COMPLEX (TRANSCRIPTION
Coumpound	NULL; 1ASU 8	INTEGRASE; CHAIN: A;	INTEGRASE; CHAIN: A;	INTEGRASE; CHAIN: A, B, C;				0 4 1 14 140	INTEGRASE; CHAIN: A, B, C,	i		RSV INTEGRASE; CHAIN: A,	B;		AVIAN SARCOMA VIRUS	INTEGRASE; CHAIN: A;	POL POLYPROTEIN; CHAIN:	A, B;		HIV-1 INTEGRASE: CHAIN:	A. B. C.				TUMOR SUPPRESSOR PIGINKAA: CHAIN: NULL:	TUMOR SUPPRESSOR	P16INK4A; CHAIN: NULL;	GA BINDING PROTEIN	ALPHA; CHAIN: A; GA	CHAIN: B. DNA: CHAIN: D. E.	T			GA BINDING PROTEIN
SeqFold score																																		72.74
PMF		0.29	0.21	0.19	<u> </u>				0.1			0.13			0.68	}	0.17			0 30	65.0				0.99	_		-						
Verify score		0.21	0.07	c	,				-0.06			-0.25	_		800	}	0.17			1	; ;				0.41	0.68		0.12						
PSI- BLAST		5.40E-26	5.40E-31	5 40E-33	J. 101-33				1.30E-29			3.60E-26			3 605 21	2.000.0	9.00E-26			00.000	9.00E-20				1.30E-24	6.50E-24		1.10E-34						5.40E-39
End		297	297	207	167				337			337			22.4	1,7	297			100	/67				228	197	<u>:</u>	239						230
· Start AA		142	142	25	74.				131			139	ì		75.	3	142				147			•	112	76	<u> </u>	112						73
Chain			*	,	ر				V			c	à		_	<	\	!			∢							В						B
PDB		1404	1691	5	FIGT		•		1c0	E		1010	1			TCXd	lexo				Iqs4				185c	1050	3	lawc						lawc
SEQ	ö	316	715	,	(1)				715			3116	}			cr/	715	:			715				719	710		719						719

					т			
PDB annotation	REGULATION/DNA) GABPALPHA; GABPBETA1; COMPLEX (TRANSCRIPTION EGULATION/DNA), DNA-BINDING, 2 NUCLEAR PROTEIN, ETS DOMAIN, ANK YRIN REPEATS, TRANSCRIPTION 3 FACTOR	COMPLEX (TRANSCRIPTION REGULATIONDNA) GABPALPHA; GABPBETA1; COMPLEX (TRANSCRIPTION REGULATIONDNA), DNA-BINDING, 2 NUCLEAR PROTEIN, ETS DOMAIN, ANKYRIN REPEATS, TRANSCRIPTION 3 FACTOR	TUMOR SUPPRESSOR TUMOR SUPPRESSOR, CDK4/6 INHIBITOR, ANK YRIN MOTIF	TUMOR SUPPRESSOR TUMOR SUPPRESSOR, CDK4/6 INHIBITOR, ANKYRIN MOTIF	TUMOR SUPPRESSOR TUMOR SUPPRESSOR, CDK4/6 INHIBITOR, ANKYRIN MOTIF	TUMOR SUPPRESSOR TUMOR SUPPRESSOR, CDK4/6 INHIBITOR, ANK YRIN MOTIF	COMPLEX (KINASE/ANTI- ONCOGENE) CDK6; P16INK44, MTS1; CYCLIN DEPENDENT KINASE, CYCLIN DEPENDENT KINASE INHIBITORY 2 PROTEIN, CDK, INK4, CELL CYCLE, MULTIPLE TUMOR SUPPRESSOR, 3 MTS1, COMPLEX (KINASE/ANTI-ONCOGENE) HEADER	COMPLEX (INHIBITOR PROTEINKINASE) INHIBITOR PROTEIN, CYCLIN-DEPENDENT KINASE, CELL CYCLE 2 CONTROL, ALPHABETA, COMPLEX (INHIBITOR PROTEIN/KINASE)
Coumpound	ALPHA; CHAIN: A: GA BINDING PROTEIN BETA I; CHAIN: B: DNA; CHAIN: D, E;	GA BINDING PROTEIN ALPHA; CHAIN: A; GA BINDING PROTEIN BETA 1; CHAIN: B; DNA; CHAIN: D, E;	P19INK4D CDK4/6 INHIBITOR; CHAIN; NULL;	P19INK4D CDK4/6 INHIBITOR; CHAIN: NULL;	P19INK4D CDK4/6 INHIBITOR; CHAIN: NULL;	P19TNK4D CDK4/6 INHIBITOR; CHAIN: NULL;	CYCLIN-DEPENDENT KINASE 6; CHAIN: A; MULTIPLE TUMOR SUPPRESSOR; CHAIN: B;	CYCLIN-DEPENDENT KINASE 6; CHAIN: A; P19INK4D; CHAIN: B;
SeqFold score		-		68.17				
PMF			0.51			-	-	0.36
Verify score		0.32	90:04		0.77	0.13	0.33	0.21
PSI- BLAST		5.40E-39	1.10E-25	7.80E-35	7.80E-35	1.10E-34	1.40E-25	9.00E-24
End		228	160	233	218	231	228	160
Start AA		78	21	73	78	78	112	21
Chain		B					ø	æ
PDB		lawc	1bd8	1bd8	1bd8	1bd8	lbi7	16lx
SEQ	Ö	719	719	719	719	719	719	719

PDB annotation	COMPLEX (INHIBITOR PROTEIN/KINASE) INHIBITOR PROTEIN/CIVASE) INHIBITOR KINASE, CELL CYCLE 2 CONTROL, ALPHA/BETA, COMPLEX (INHIBITOR PROTEIN/KINASE)	COMPLEX (INHIBITOR PROTEIN/KINASE) INHIBITOR PROTEIN/CYCLIA-DEPENDENT KINASE, CELL CYCLE 2 CONTROL, ALPHA/BETA, COMPLEX (INHIBITOR PROTEIN/KINASE)	COMPLEX (INHIBITOR PROTEIN/KINASE) INHIBITOR PROTEIN, CYCLIN-DEPENDENT KINASE, CELL CYCLE 2 CONTROL, ALPHA/BETA, COMPLEX (INHIBITOR PROTEIN/KINASE)	HORMONE/GROWTH FACTOR P18- INK4C; CELL CYCLE INHIBITOR, P18INK4C, TUMOR, SUPPRESSOR. CYCLIN- 2 DEPENDENT KINASE, HORMONE/GROWTH FACTOR	HORMONE/GROWTH FACTOR P18- INK4C, CELL CYCLE INHIBITOR, P18INK4C, TUMOR, SUPPRESSOR, CYCLIN- 2 DEPENDENT KINASE, HORMONE/GROWTH FACTOR	HORMONEGROWTH FACTOR P18- INK4C; CELL CYCLE INHIBITOR, P18INK4C; TUMOR, SUPPRESSOR, CYCLIN-2 DEPENDENT KINASE, HORMONE/GROWTH FACTOR	SIGNALING PROTEIN HELIX-TURN- HELIX, ANKYRIN REPEAT	SIGNALING PROTEIN HELIX-TURN- HELIX, ANKYRIN REPEAT	SIGNALING PROTEIN HELIX-TURN- HELIX, ANKYRIN REPEAT
Coumpound	CYCLIN-DEPENDENT KINASE 6; CHAIN: A; P19INK4D; CHAIN: B;	CYCLIN-DEPENDENT KINASE 6: CHAIN: A; P19INK4D; CHAIN: B;	CYCLIN-DEPENDENT KINASE 6; CHAIN: A; P19INK4D; CHAIN: B;	CYCLIN-DEPENDENT KINASE 6 INHIBITOR; CHAIN: A;	CYCLIN-DEPENDENT KINASE 6 INHIBITOR; CHAIN: A;	CYCLIN-DEPENDENT KINASE 6 INHIBITOR; CHAIN: A;	CYCLIN-DEPENDENT KINASE 4 INHIBITOR B; CHAIN: A;	CYCLIN-DEPENDENT KINASE 4 INHIBITOR B; CHAIN: A;	CYCLIN-DEPENDENT KINASE 4 INHIBITOR B;
SeqFold score	70.26				67.41				
PMF		-	-	0.12			-	0.99	_
Verify score		0.62	0.39	-0.07		0.17	0.33	0.48	0.73
PSI- BLAST	1.30E-35	1.30E-35	3.60E-34	1.10E-25	1.105-36	1.10E-36	2.60E-29	1.10E-25	1.20E-28
End	201	218	231	165	239	233	218	234	197
Start AA	44	78	84	18	70	78	101	112	78
Chain	æ	В	B	∢	∢	4	¥_	4	4
PDB	16lx	1blx	. Ibix	1bu9	1bu9	1bu9	149s	1d9s	1d9s
SEQ NO.	719	719	719	719	719	719	719	719	719

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PDB annotation	C.4 (C)	CELL CYCLE INHIBITOR PT8- INKAC(INK6); CELL CYCLE INHIBITOR, P18-INKAC(INK6), ANKYRIN REPEAT, 2 CDK 4/6 INHIBITOR	CELL CYCLE INHIBITOR P18- INK4C(INK6); CELL CYCLE INHIBITOR, P18-INK4C(INK6), ANKYRIN REPEAT, 2 CDK 4/6 INHIBITOR F PARTICULATION P18	CELL CYCLE INHIBITOR F18- INKAC(INK6); CELL CYCLE INHIBITOR, F18-INKAC(INK6), ANKYRIN REPEAT, 2 CDK 4/6 INHIBITOR	CELL CYCLE INHIBITOR P18- INKAC(INK6); CELL CYCLE INHIBITOR, P18-INKAC(INK6), ANK YRIN REPEAT, 2 CDK 4/6 INHIBITOR	TRANSCRIPTION FACTOR P65; P30D; TRANSCRIPTION FACTOR, IKBNFKB COMPLEX	TRANSCRIPTION FACTOR P65; P30D; TRANSCRIPTION FACTOR, IKBNFKB COMPLEX	TRANSCRIPTION FACTOR Post Pob; TRANSCRIPTION FACTOR, IKBAFKB COMPLEX	TRANSCRIPTION FACTOR P65; P50D; TRANSCRIPTION FACTOR, IKBNFKB COMPLEX	TRANSCRIPTION FACTOR P65; P50D; TRANSCRIPTION FACTOR, IKBNFKB COMPLEX
Commonud	CHAIN: A;	CYCLIN-DEPENDENT KINASE 6 INHIBITOR; CHAIN: A, B;	CYCLIN-DEPENDENT KINASE 6 INHIBITOR; CHAIN: A, B;	CYCLIN-DEPENDENT KINASE 6 INHIBITOR; CHAIN: A, B;	CYCLIN-DEPENDENT KINASE 6 INHIBITOR; CHAIN: A, B;	NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF-KAPPA-B P50D SUBUNIT; CHAIN: C; I- KAPPA-B-ALPHA; CHAIN: D;	NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF-KAPPA-B PSOD SUBUNIT; CHAIN: C; I- KAPPA-B-ALPHA; CHAIN: D;	NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF-KAPPA-B P50D SUBUNIT; CHAIN: C; I- KAPPA-B-ALPHA; CHAIN: D;	NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF-KAPPA-B PS0D SUBUNIT: CHAIN: C; I- KAPPA-B-ALPHA; CHAIN: D;	NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF-KAPPA-B P50D SUBUNIT; CHAIN: C; I- KAPPA-B-ALPHA; CHAIN: D;
SeqFold score				63.76				69.17		
PMF		90.0	0.93		-	0.33	6.0		69:0	0.76
Verify score		-0.2	0.31		0.24	-0.33	-0.04		-0.05	-0.21
PSI- BLAST		1.40E-21	5.40E-25	5.40E-36	5.40E-36	3.60E-26	3.60E-33	7.20E-39	7.20E-39	1.30E-27
End		131	164	232	232	238	177	216	228	144
Start		-	81	75	78	107	13	21	39	8
Chain 10		<	<	A	<	Ω	Ω	D	Q	Q
PDB D	1	lihb	lihb	di:I	1ihb	1 ika	likn	likn	lika Pika	likn
SEQ	ğ	719	719	719	719	719	719	.612	719	617

PDB annotation	ANK-REPEAT MYOTROPHIN, ACETYLATION, NMR, ANK-REPEAT	ANK-REPEAT MYOTROPHIN, ACETYLATION, NMR, ANK-REPEAT	COMPLEX (KANSCERFIUN REGANK REPEAT) COMPLEX (TRANSCRIPTION REGULATION/ANK REPEAT), ANKYRIN 2 REPEAT HELIX	COMPLEX (KANSCRIPTION REGANK REPEAT) COMPLEX (TRANSCRIPTION REGULATION/ANK REPEAT), ANKYRINZ REPEAT HELIX	COMPLEX (IRANSCIPTION COMPLEX (TRANSCIPTION REGULATION/ANK REPEAT), ANKYRIN 2 REFEAT HELIX	COMPLEX (TRANSCRIPTION REG/ANK REPEAT) COMPLEX (TRANSCRIPTION REGULATION/ANK REPEAT), ANKYRIN 2 REPEAT HELIX	COMPLEX (ANTI- ONCOGENE/ANKYRIN REPEATS) P33BP2; ANKYRIN REPEATS, SH3, P53, TUMOR SUPPRESSOR, MULTIGENE 2 FAMILY, NUCLEAR PROTEIN, PHOSPHORYLATION, DISEASE MUTATION, 3 POLYMORPHISM, COMPLEX (ANTI- ONCOGENE/ANTI-	COMPLEX (TRANSCRIPTION REGULATION/DNA) GABPALPHA; GABPBETAI; COMPLEX (TRANSCRIPTION REGULATION/DNA), DNA-BINDING, 2 NUCLEAR PROTEIN, ETS DOMAIN, ANKYRIN REPEATS, TRANSCRIPTION 3 FACTOR	COMPLEX (TRANSCRIPTION REGULATION/DNA) GABPALPHA; GABPBETAI; COMPLEX
Coumpound	MYOTROPHIN; CHAIN: NULL	MYOTROPHIN; CHAIN: NULL	NF-KAPPA-B P65; CHAIN: A, C; NF-KAPPA-B P50; CHAIN: B, D; I-KAPPA-B-ALPHA; CHAIN: E, F;	NF-KAPPA-B P65; CHAIN: A, C; NF-KAPPA-B P50; CHAIN: B, D; I-KAPPA-B-ALPHA; CHAIN: E, F;	NF-KAPPA-B P65; CHAIN: A, C; NF-KAPPA-B P50; CHAIN: B, D; I-KAPPA-B-ALPHA; CHAIN: E, F;	NF-KAPPA-B P65; CHAIN: A, C; NF-KAPPA-B P50; CHAIN: B, D; I-KAPPA-B-ALPHA; CHAIN: E, F;	P53; CHAIN: A; 53BP2; CHAIN: B;	GA BINDING PROTEIN ALPHA; CHAIN: A; GA BINDING PROTEIN BETA 1; CHAIN: B; DNA; CHAIN: D, E;	GA BINDING PROTEIN ALPHA; CHAIN: A; GA BINDING PROTEIN BETA I;
SeqFold		64.15				65.66	64.29		
PMF			0.94	0.92	0.78			1	-
Verify	0.14		0.19	-0.07	0.07			0.79	0.83
PSI- BLAST	2.60E-28	6.50E-33	9.00E-26	9.00E-33	1.10E-38	1.10E-38	1.60E-20	7.80E-42	2.60E-42
End	225	192	238	177	228	202	239	312	147
Start	110	75	106	12	39	7	74	161	2.
Chain			យ	ш	E	ъ	Ø	<u>м</u>	В
PDB D	1my	1my		lnfi	Jufi	Infi	1ycs	lawc	1awc
<u> </u>	NO.	719	719	719	719	719	719	721	721

PDB annotation	(TRANSCRIPTION REGULATION/DNA), DNA-BINDING, 2 NUCLEAR PROTEIN, ETS DOMAIN, ANKYRIN REPEATS, TRANSCRIPTION 3 FACTOR	COMPLEX (TRANSCRIPTION RECULATION/DNA) GABPALPHA; GABPBETAI; COMPLEX GRANSCRIPTION RECULATION/DNA), DNA-BINDING, 2 RECULATION/DNA), DNA-BINDING, 2 NUCLEAR PROTEIN, ETS DOMAIN, ANKYRIN REPEATS, TRANSCRIPTION 3 FACTOR	COMPLEX (TRANSCRIPTION REGULATION/DNA) GABPALPHA; GABPBETA1; COMPLEX (TRANSCRIPTION REGULATION/DNA), DNA-BINDING, 2 NUCLEAR PROTEIN, ETS DOMAIN, ANKYRIN REPEATS, TRANSCRIPTION 3 FACTOR	COMPLEX (TRANSCRIPTION REGULATIONDNA) GABPALPHA; GABPBETA; COMPLEX (TRANSCRIPTION REGULATIONDNA), DNA-BINDING, 2 NUCLEAR PROTEIN, ETS DOMAIN, ANKYRIN REPEATS, TRANSCRIPTION 3 FACTOR	COMPLEX (TRANSCRIPTION REGULATION/DNA) GABPALPHA; GABPBETA1; COMPLEX GRANSCRIPTION REGULATION/DNA, DNA-BINDING, 2 NUCLEAR PROTEIN, ETS DOMAIN, ANKYRIN REPEATS, TRANSCRIPTION 3 FACTOR	COMPLEX (TRANSCRIPTION REGULATION/DNA) GABPALPHA; GABPBETA1; COMPLEX (TRANSCRIPTION
Coumpound	CHAIN: B; DNA; CHAIN: D, E;	GA BINDING PROTEIN ALPHA; CHAIN: A; GA BINDING PROTEIN BETA 1; CHAIN: B; DNA; CHAIN: D, E;	GA BINDING PROTEIN ALPHA; CHAIN: A; GA BINDING PROTEIN BETA 1; CHAIN: B; DNA; CHAIN: D, E;	GA BINDING PROTEIN ALPHA; CHAIN: A; GA BINDING PROTEIN BETA 1; CHAIN: B; DNA; CHAIN: D, E;	GA BINDING PROTEIN ALPHA; CHAIN: A; GA BINDING PROTEIN BETA 1; CHAIN: B; DNA; CHAIN: D, E;	GA BINDING PROTEIN ALPHA; CHAIN: A; GA BINDING PROTEIN BETA 1; CHAIN: B; DNA; CHAIN: D, E;
SeqFold score						
PMF score			<u>.</u>			-
Verify		0.71	1.14	0.72	6.0	1.23
PSI- BLAST		3.60E-34	5.20E-43	1.30E-44	3.60E-39	5.20E-45
End		147	378	180	378	412
Start AA		2	226		231	259
Chain		æ	m .	æ	ø	æ
PDB ID		lawc	lawc	lawc	lawc	lawc
SEQ	Ö	721	721	721	721	721

						
PDB annotation	REGULATION/DNA), DNA-BINDING, 2 NUCLEAR PROTEIN, ETS DOMAIN, ANKYRIN REPEATS, TRANSCRIPTION 3 FACTOR	COMPLEX (TRANSCRIPTION REGULATION/DNA) GABPALPHA; GABPBETA1; COMPLEX (TRANSCRIPTION REGULATION/DNA, DNA-BINDING, 2 NUCLEAR PROTEIN, ETS DOMAIN, ANKYRIN REPEATS, TRANSCRIPTION 3 FACTOR	COMPLEX (TRANSCRIPTION REGULATION/DNA) GABPALPHA; GABPBETA1; COMPLEX (TRANSCRIPTION REGULATION/DNA, DNA-BINDING, 2 NUCLEAR PROTEIN, ETS DOMAIN, ANK YRIN REPEATS, TRANSCRIPTION 3 FACTOR	COMPLEX (TRANSCRIPTION REGULATIONDNA) GABPALPHA; GABPBETA1; COMPLEX (TRANSCRIPTION REGULATIONDNA), DNA-BINDING. 2 NUCLEAR PROTEIN, ETS DOMAIN, ANKYRIN REPEATS, TRANSCRIPTION 3 FACTOR	COMPLEX (TRANSCRIPTION REGULATIONDNA) GABPALPHA; GABPBETAI; COMPLEX (TRANSCRIPTION REGULATIONDNA), DNA-BINDING, 2 NUCLEAR PROTEIN, ETS DOMAIN, ANKYRIN REPEATS, TRANSCRIPTION 3 FACTOR.	COMPLEX (TRANSCRIPTION REGULATION/DNA) GABPALPHA; GABPBETA1; COMPLEX (TRANSCRIPTION REGULATION/DNA), DNA-BINDING, 2
Coumpound		GA BINDING PROTEIN ALPHA; CHAIN: A; GA BINDING PROTEIN BETA 1; CHAIN: B; DNA; CHAIN: D, E;	GA BINDING PROTEIN ALPHA; CHAIN: A; GA BINDING PROTEIN BETA I; CHAIN: B; DNA; CHAIN: D; E;	GA BINDING PROTEIN ALPHA; CHAIN: A: GA BINDING PROTEIN BETA I; CHAIN: B: DNA; CHAIN: D, E;	GA BINDING PROTEIN ALPHA; CHAIN: A; GA BINDING PROTEIN BETA 1; CHAIN: B; DNA; CHAIN: D, E;	GA BINDING PROTEIN ALPHA; CHAIN: A; GA BINDING PROTEIN BETA 1; CHAIN: B; DNA; CHAIN: D, E;
SeqFold score						
PMF			•	-		0.62
Verify score		0.93	0.93	0.69		0.22
PSI- BLAST		5.40E-41	1.80E-36	1.30E-38	1.80E-36	7.20E-32
End		411	442	180	475	487
Start		264	297	33		363
Chain		В	В	ø	α .	В
PDB ID		lawc	lawc	lawc	lawc	lawc
SEQ	Ö	721	721	721	721	721

PDB annotation	NUCLEAR PROTEIN, ETS DOMAIN, ANKYRIN REPEATS, TRANSCRIPTION 3 FACTOR	COMPLEX (TRANSCRPTION REGULATION/DNA) GABPALPHA; GABPBETA; COMPLEX (TRANSCRIPTION REGULATION/DNA), DNA-BINDING, 2 NUCLEAR PROTEIN, ETS DOMAIN, ANKYRIN REPEATS, TRANSCRIPTION 3 FACTOR	COMPLEX (IKANSCRIFILD) REGULATION/DNA) GABPBETA1; COMPLEX (TRANSCRIPTION REGULATION/DNA), DNA-BINDING, 2 NUCLEAR PROTEIN, ETS DOMAIN, ANKYRIN REPEATS, TRANSCRIPTION 3 FACTOR	COMPLEX (TRANSCRIPTION REGULATIONDNA) GABPALPHA; GABPBETAI; COMPLEX (TRANSCRIPTION REGULATIONDNA), DNA-BINDING, 2 NUCLEAR PROTEIN, ETS DOMAIN, ANKYRIN REPEATS, TRANSCRIPTION 3 FACTOR	TUMOR SUPPRESSOR TUMOR SUPPRESSOR, CDK4/6 INHIBITOR, ANKYRIN MOTIP	TUMOR SUPPRESSOR TUMOR SUPPRESSOR, CDK4/6 INHIBITOR, ANKYRIN MOTIF	TUMOR SUPPRESSOR TUMOR SUPPRESSOR, CDK4/6 INHIBITOR, ANKYRIN MOTIF	TUMOR SUPPRESSOR TUMOR SUPPRESSOR, CDK4/6 INHIBITOR, ANK YRIN MOTIF	TUMOR SUPPRESSOR TUMOR SUPPRESSOR, CDK4/6 INHIBITOR,
Coumpound		GA BINDING PROTEIN ALPHA; CHAIN: A; GA BINDING PROTEIN BETA I; CHAIN: B; DNA; CHAIN: D, E;	GA BINDING PROTEIN ALPHA; CHAIN: A; GA BINDING PROTEIN BETA 1; CHAIN: B; DNA; CHAIN: D, E;	GA BINDING PROTEIN ALPHA; CHAIN: A; GA BINDING PROTEIN BETA 1; CHAIN: B; DNA; CHAIN: D, E;	P19TNK4D CDK4/6 INHIBITOR; CHAIN: NULL;	P19INK4D CDK4/6 INHIBITOR; CHAIN: NULL;	P19INK4D CDK4/6 INHIBITOR; CHAIN: NULL;	P19INK4D CDK4/6 INHIBITOR; CHAIN: NULL;	P19TNK4D CDK4/6 INHIBITOR; CHAIN: NULL;
SeqFold score									
PMF		1	-	-	-	0.42	-		-
Verify score		0.59	0.79	0.87	0.79	0.12	0.43	0.73	0.53
PSI- BLAST		6.50E.47	1.40E-39	5.40E-38	2.60E-38	2.60E-38	1.20E-40	9.10E-41	1.30E-41
End		213	213	246	281	348	149	381	182
Start AA		61	99	66	128	161	7	227	24
Chain		m	æ	<u>α</u>					
PDB		lawc	lawc	lawc .	1bd8	1bd8	1bd8	1bd8	1 pd8
SEQ	O	721	721	721	721	721	721	121	721

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PDB annotation	ANKYRIN MOTIF	TUMOR SUPPRESSOR TUMOR SUPPRESSOR, CDK4/6 INHIBITOR, ANK YRIN MOTIF	TUMOR SUPPRESSOR TUMOR SUPPRESSOR, CDK4/6 INHIBITOR, ANK YRIN MOTIF	TUMOR SUPPRESSOR TUMOR SUPPRESSOR, CDK4/6 INHIBITOR, ANK YRIN MOTIF	COMPLEX (INHIBITOR PROTEIN/KINASE) INHIBITOR PROTEIN, CYCLIN-DEPENDENT	KINASE, CELL CYCLE 2 CONTROL, ALPHA/BETA, COMPLEX (INHIBITOR PROTEIN/KINASE)	COMPLEX (INHIBITOR PROTEIN/KINASE) INHIBITOR	PROTEIN, CYCLIN-DEPENDENT KINASE, CELL CYCLE 2 CONTROL,	ALPHA/BETA, COMPLEX (INHIBITOR PROTEIN/KINASE)	COMPLEX (INHIBITOR	PROTEIN/KINASE) INHIBITOR	KINASE, CELL CYCLE 2 CONTROL,	ALPHA/BETA, COMPLEX (INHIBITOR PROTEIN/KINASE)	COMPLEX (INHIBITOR	PROTEIN/KINASE) INHIBITOR PROTEIN, CYCLIN-DEPENDENT	KINASE, CELL CYCLE 2 CONTROL,	ALPHA/BETA, COMPLEX (INHIBITOR PROTEIN/KINASE)	COMPLEX (INHIBITOR	PROTEIN/KINASE) INHIBITOR	KINASE, CELL CYCLE 2 CONTROL,	ALPHA/BETA, COMPLEX (INHIBITOR PROTEIN/KINASE)	COMPLEX (INHIBITOR
Coumpound		P19INK4D CDK4/6 INHIBITOR; CHAIN: NULL;	PI9NK4D CDK4/6 INHIBITOR; CHAIN: NULL;	P19INK4D CDK4/6 INHIBITOR; CHAIN: NULL;	CYCLIN-DEPENDENT KINASE 6; CHAIN: A; P19INK4D: CHAIN: B:		CYCLIN-DEPENDENT KINASE 6; CHAIN: A;	P19INK4D; CHAIN: B;		CYCLIN-DEPENDENT	KINASE 6; CHAIN: A;	PISINK4U; CHAIN: B;		CYCLIN-DEPENDENT	KINASE 6; CHAIN: A; P19INK4D: CHAIN: B:			CYCLIN-DEPENDENT	KINASE 6; CHAIN: A;	i Division, Circuit. D.		CYCLIN-DEPENDENT
SeqFold																						
PMF		~	_		-		-1		•	_				_				_				-
Verify score		0.89	0.74	29.0	0.83		0.13			0.81				0.61				0.78				1.13
PSI- BLAST		1.20E-38	1.40E-30	7.80E-44	1.30E-39		5.20E-38			1.00E-41				6.50E-43	-			1.30E-43				5.20E-41
End		414	442	215	285		350			153				185				385				416
Start AA		263	297	79	132		163			2				22				230				263
Chain					м		В			В				В				В				В
PDB		1bd8	8pq1	1bd8	16k		1blx			1bk				1blx				1blx				1blx
SEQ B B B		721	721	721	121		721			721				721				721				721

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PDB annotation	PROTEINKINASE) INHIBITOR PROTEIN, CYCLIN-DEPENDENT KINASE, CELL CYCLE 2 CONTROL, ALPHABETA, COMPLEX (INHIBITOR PROTEINKINASE)	COMPLEX (INHIBITOR PROTEIN/KINASE) INHIBITOR PROTEIN, CYCLIN-DEPENDENT KINASE, CELL CYCLE 2 CONTROL, ALPHABETA, COMPLEX (INHIBITOR PROTEIN/KINASE)	HORMONEGROW IN FACTOR TO MY 45 CELL CYCLE INHIBITOR, PI 81NK4C, TUMOR, SUPPRESSOR, CYCLIN- 2 DEPENDENT KINASE, HORMONE/GROWTH FACTOR	HORMONEGROWTH FACTOR P18- NK4C; CELL CYCLE INHIBITOR, P18INK4C, TUMOR, SUPPRESSOR, CYCLIN-2 DEPENDENT KINASE, HORMONE/GROWTH FACTOR	HORMONEGROWTH FACTOR P18- MRAG; CELL CYCLE INHIBITOR, P18INKAC, TUMOR, SUPPRESSOR, CYCLIN- 2 DEPENDENT KINASE, HORMONE/GROWTH FACTOR	SIGNALING PROTEIN HELIX-TURN- HELIX, ANKYRIN REPEAT	SIGNALING PROTEIN HELIX-TURN- HELIX, ANK YRIN REPEAT	SIGNALING PROTEIN HELIX-TURN- HELIX, ANKYRIN REPEAT	SIGNALING PROTEIN HELIX-TURN- HELIX, ANKYRIN REPEAT	SIGNALING PROTEIN HELIX-TURN- HELIX, ANKYRIN REPEAT
Coumpound	KINASE 6; CHAIN: A; P19INK4D; CHAIN: B;	CYCLIN-DEPENDENT KINASE 6; CHAIN: A; PI9INK4D; CHAIN: B;	CYCLIN-DEPENDENT KINASE 6 INHIBITOR; CHAIN: A;	CYCLIN-DEPENDENT KINASE 6 INHIBITOR; CHAIN: A;	CYCLIN-DEPENDENT KINASE 6 INHIBITOR; CHAIN: A;	CYCLIN-DEPENDENT KINASE 4 INHIBITOR B; CHAIN: A;	CYCLIN-DEPENDENT KINASE 4 INHIBITOR B; CHAIN: A;	CYCLIN-DEPENDENT KINASE 4 INHIBITOR B; CHAIN: A;	CYCLIN-DEPENDENT KINASE 4 INHIBITOR B; CHAIN: A;	CYCLIN-DEPENDENT KINASE 4 INHIBITOR B; CHAIN: A;
SeqFold score					90.26					
PMF		-	1	0.35			-		_	
Verify score		0.39	0.61	0.37		9.0	99.0	0.58	0.65	0.61
PSI- BLAST		6.50E-43	7.20E-38	1.80E-32	7.20E-38	6.50E-36	1.30E-37	1.30E-40	2.60E-38	2.60E-38
End		252	185	480	258	153	384	416	185	216
Start		64	33	330	16	14	250	283	52	84
Chain		æ	4	∢	∢	∢	4	4	4	4
PDB UD		1blx	1bu9	1bu9	16u9	1d9s	1d9s	1d9s	149s	1d9s
SEQ	Ö	721	721	721	121	721	721	721	721	121

PDB annotation	CELL CYCLE INHIBITOR P18- INK4C(INK6); CELL CYCLE INHIBITOR, P18-INK4C(INK6), ANKYRIN REPEAT, 2 CDK 4/6 INHIBITOR	CELL CYCLE INHIBITOR P18- INK4C(INK6); CELL CYCLE INHIBITOR, P18-INK4C(INK6), ANKYRIN REPEAT, 2 CDK 4/6 INHIBITOR	CELL CYCLE INHIBITOR P18- INK4C(INK6); CELL CYCLE INHIBITOR, P18-INK4C(INK6), INKYRIN REPEAT, 2 CDK 4/6 INHIBITOR	CELL CYCLE INHIBITOR P. 18- INK4C(INK6); CELL CYCLE INHIBITOR, P. 18-INK4C(INK6), ANK YRIN REPEAT, 2 CDK 4/6 INHIBITOR	TRANSCRIPTION FACTOR P65, P50D; TRANSCRIPTION FACTOR, IKBNFKB COMPLEX	TRANSCRIPTION FACTOR P65; P50D; TRANSCRIPTION FACTOR, IKBNFKB COMPLEX	TRANSCRIPTION FACTOR P65; P50D; TRANSCRIPTION FACTOR, IKBNFKB COMPLEX	TRANSCRIPTION FACTOR P65; P50D; TRANSCRIPTION FACTOR, IKBANFKB COMPLEX	TRANSCRIPTION FACTOR P65; P50D; TRANSCRIPTION FACTOR, IKBANFKB COMPLEX	TRANSCRIPTION FACTOR P65; P50D;
Coumpound			CYCLIN-DEPENDENT CEL. KINASE 6 INHIBITOR; INK. CHAIN: A, B; ANK. ANKI	CYCLIN-DEPENDENT CEL. KINASE 6 INHIBITOR; INK CHAIN: A, B; ANK ANK INH	NF-KAPPA-B P65 SUBUNIT; TRA CHAIN: A; NF-KAPPA-B P50D TRA SUBUNIT; CHAIN: C; 1- KAPPA-B-ALPHA; CHAIN: D;	NF-KAPPA-B P65 SUBUNIT; TRA CHAIN: A; NF-KAPPA-B P50D TRA SUBUNIT; CHAIN: C; I- KAPPA-B-ALPHA; CHAIN: D;		— .;		Н
SeqFold score				88.22	82.11					
PMF		-	0.92			0.93	66:0	-	1	0.39
Verify score	0.7	0.55	0.5			-0.02	0.48	0.56	0.55	0.4
PSI- BLAST	1.80E-32	3.60E-37	1.30E-31	3.60E-37	2.60E-57	2.60E-55	5.20E-52	5.40E-43	7.80E-52	7.20E-39
End	446	184	479	250	333	386	188	411	416	475
Start	297	33	330	96	127	161	7	226	226	292
Chain ID	¥.	V	4	< _	Ω	Ω	Ω	Q	О	Д
PDB	lihb	1ihb	lihb	lihb	lika	lika	1 ika	1 ika	1 iku	likn
SEQ D	721	721	721	121	721	721	721	721	721	721

											
PDB annotation	TRANSCRIPTION FACTOR, IKBNFKB COMPLEX	TRANSCRIPTION FACTOR P65; P50D; TRANSCRIPTION FACTOR, IKBNFKB COMPLEX	TRANSCRIPTION FACTOR P65; P50D; TRANSCRIPTION FACTOR, IKB/NFKB COMPLEX	TRANSCRIPTION FACTOR P65, P50D; TRANSCRIPTION FACTOR, IKBNFKB COMPLEX	COMPLEX (TRANSCRIPTION REG/ANK REPEAT) COMPLEX (TRANSCRIPTION REGULATION/ANK REPEAT), ANKYRIN 2 REPEAT HELIX	COMPLEX (TRANSCRIPTION REG/ANK REPEAT) COMPLEX (TRANSCRIPTION REGULATION/ANK REPEAT), ANKYRIN 2 REPEAT HELIX	COMPLEX (TRANSCRIPTION REG/ANK REPEAT) COMPLEX (TRANSCRIPTION REGULATION/ANK REPEAT), ANKYRIN 2 REPEAT HELIX	COMPLEX (TRANSCRIPTION REG/ANK REPEAT) COMPLEX (TRANSCRIPTION REGULATION/ANK REPEAT), ANKYRIN 2 REPEAT HELIX	COMPLEX (TRANSCRIPTION REG/ANK REPEAT) COMPLEX (TRANSCRIPTION REGULATION/ANK REPEAT), ANKYRIN 2 REPEAT HELIX	COMPLEX (TRANSCRIPTION REG/ANK REPEAT) COMPLEX (TRANSCRIPTION REGULATION/ANK REPEAT), ANKYRIN 2 REPEAT HELIX	COMPLEX (TRANSCRIPTION REG/ANK REPEAT) COMPLEX
Coumpound	CHAIN: A; NF-KAPPA-B P50D SUBUNIT; CHAIN: C; I- KAPPA-B-ALPHA; CHAIN: D;	NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NP-KAPPA-B P50D SUBUNIT; CHAIN: C; 1- KAPPA-B-ALPHA; CHAIN: D;	NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NP-KAPPA-B P50D SUBUNIT; CHAIN: C; I- KAPPA-B-ALPHA; CHAIN: D;	NF.KAPPA-B P65 SUBUNIT; CHAIN: A; NF.KAPPA-B P50D SUBUNIT; CHAIN: C; I- KAPPA-B-ALPHA; CHAIN: D;	NF-KAPPA-B P65; CHAIN: A, C; NF-KAPPA-B P50; CHAIN: B, D; I-KAPPA-B-ALPHA; CHAIN: E, F;	NF-KAPPA-B P65; CHAIN: A, C; NF-KAPPA-B P50; CHAIN: B, D; I-KAPPA-B-ALPHA; CHAIN: E, F;	NF-KAPPA-B P65; CHAIN: A, C; NF-KAPPA-B P50; CHAIN: B, D; I-KAPPA-B-ALPHA; CHAIN: E, F;	NF-KAPPA-B P65; CHAIN: A, C; NF-KAPPA-B P50; CHAIN: B, D; I-KAPPA-B-ALPHA; CHAIN: E, F;	NF-KAPPA-B P65; CHAIN: A, C; NF-KAPPA-B P50; CHAIN: B, D; L-KAPPA-B-ALPHA; CHAIN: E, F;	NF-KAPPA-B P65; CHAIN: A, C; NF-KAPPA-B P50; CHAIN: B, D; I-KAPPA-B-ALPHA; CHAIN: E, F;	NF-KAPPA-B P65; CHAIN: A, C; NF-KAPPA-B P50; CHAIN:
SeqFold score				,	85.42						
PMF		0.05	-	1		,	1	1	-	_	_
Verify score		0.09	0.2	0.22		0.08	69:0	0.49	0.82	8.0	0.65
PSI- BLAST		7.20E-34	5.40E-38	2.60E-57	2.60E-52	2.60E-52	7.80E-42	7.80E-53	7.20E-43	6.50E-52	1.60E-38
End		486	233	256	322	350	153	216	411	420	475
Start		325	19	61	124	124	7	21	224	226	292
Chain		Ω	Q	D	a	ല	ш	ы	ല	മ	ы
PDB		likn	likn	n lika	Infi	Infi	lnfi	1nfi	1nfi	1nfi	Inf
SEQ EQ		721	721	721	721	721	721	721	721	721	121

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PDB annotation	(TRANSCRIPTION REGULATION/ANK REPEAT), ANKYRIN 2 REPEAT HELIX	COMPLEX (TRANSCRIPTION REG/ANK REPEAT) COMPLEX (TRANSCRIPTION REGULATION/ANK REPEAT), ANKYRIN 2 REPEAT HELIX	COMPLEX (TRANSCRIPTION REGULATION/DIA) GABPALPHA; GABPBETA1; COMPLEX GRANSCRIPTION REGULATION/DIA), DNA-BINDING, 2 NUCLEAR PROTEIN, ETS DOMAIN, ANKYRIN REPEATS, TRANSCRIPTION 3 FACTOR	COMPLEX (TRANSCRIPTION REGULATION/DNA) GABPALPHA; GABPBETA1; COMPLEX (TRANSCRIPTION REGULATION/DNA), DNA-BINDING, 2 NUCLEAR PROTEIN, ETS DOMAIN, ANKYRIN REPEATS, TRANSCRIPTION 3 FACTOR	COMPLEX (TRANSCRIPTION REGULATION/DINA) GABPALPHA; GABPBETA1; COMPLEX (TRANSCRIPTION REGULATION/DINA), DINA-BINDING, 2 NUCLEAR PROTEIN, ETS DOMAIN, ANKYRIN REPEATS, TRANSCRIPTION 3 FACTOR	COMPLEX (TRANSCRIPTION REGULATION/DNA) GABPALPHA; GABPBETA1; COMPLEX (TRANSCRIPTION (TRANSCRIPTION) TOUCLEAR PROTEIN, ETS DOMAIN, ANKYRIN REPEATS, TRANSCRIPTION 3 FACTOR	COMPLEX (TRANSCRIPTION REGULATION/DNA) GABPALPHA;
Coumpound	B, D; I-KAPPA-B-ALPHA; CHAIN: E, F;	NF-KAPPA-B P65; CHAIN: A, C; NF-KAPPA-B P50; CHAIN: B, D; I-KAPPA-B-ALPHA; CHAIN: E, F;	GA BINDING PROTEIN ALPHA; CHAIN: A; GA BINDING PROTEIN BETA 1; CHAIN: B; DNA; CHAIN: D, E;	GA BINDING PROTEIN ALPHA: CHAIN: A; GA BINDING PROTEIN BETA 1; CHAIN: B; DNA; CHAIN: D, E;	GA BINDING PROTEIN ALPHA; CHAIN: A; GA BINDING PROTEIN BETA 1; CHAIN: B; DNA; CHAIN: D, E;	GA BINDING PROTEIN ALPHA; CHAIN: A; GA BINDING PROTEIN BETA 1; CHAIN: B; DNA; CHAIN: D, E;	GA BINDING PROTEIN ALPHA; CHAIN: A; GA
SeqFold score						64.41	
PMF		96.0		0.64	-0.17		-
Verify score		0.52	0.32	0.4	0.02		0.35
PSI- BLAST		1.80E-33	1.60E-31	1.60E-23	3.60E-30	5.40E-36	5.40E-36
End		486	280	343	408	198	157
Start		323	107	175	265	31	9
Chain TD		阳	В	æ	æ	m .	В
PDB		lnfi	lawc	lawc	lawc	lawc	lawc
SEQ B	Ë	721	723	723	723	723	723

PDB annotation	GABPBETA1; COMPLEX (TRANSCRIPTION REGULATIONDNA), DNA-BINDING, 2 NUCLEAR PROTEIN, ETS DOMAIN, 3 FACTOR	COMPLEX (TRANSCRIPTION REGULATION/DNA) GABPALPHA; GABPBETA1; COMPLEX (TRANSCRIPTION REGULATION/DNA), DNA-BINDING, 2 NUCLEAR PROTEIN, ETS DOMAIN, ANK YRIN REPEATS, TRANSCRIPTION 3 FACTOR	TUMOR SUPPRESSOR TUMOR SUPPRESSOR, CDK4/6 INHIBITOR, ANK YRIN MOTIF	TUMOR SUPPRESSOR TUMOR SUPPRESSOR, CDK4/6 INHIBITOR, ANKYRIN MOTIF	TUMOR SUPPRESSOR TUMOR SUPPRESSOR, CDK4/6 INHIBITOR, ANKYRIN MOTIF	COMPLEX (INHIBITOR PROTEIN/KINASE) INHIBITOR PROTEIN, CYCLIN-DEPENDENT KINASE, CELL CYCLE 2 CONTROL, ALPHA/BETA, COMPLEX (INHIBITOR PROTEIN/KINASE)	COMPLEX (INHIBITOR PROTEIN/KINASE) INHIBITOR PROTEIN, CYCLIM-DEPENDENT KINASE, CELL CYCLE 2 CONTROL, ALPHABETA, COMPLEX (INHIBITOR PROTEIN/KINASE)	HORMONE/GROWTH FACTOR P18- INK4C; CELL CYCLE INHIBITOR, P18INK4C; TUMOR, SUPPRESSOR, CYCLIN-2 DEPENDENT KINASE, HORMONE/GROWTH FACTOR	HORMONE/GROWTH FACTOR P18-
Coumpound	BINDING PROTEIN BETA 1; CHAIN: B; DNA; CHAIN: D, E;	GA BINDING PROTEIN ALPHA; CHAIN: A; GA BINDING PROTEIN BETA 1; CHAIN: B; DNA; CHAIN: D, E:	P19NK4D CDK4/6 INHIBITOR; CHAIN: NULL;	P19INK4D CDK4/6 INHIBITOR; CHAIN: NULL;	P19INK4D CDK4/6 INHIBITOR; CHAIN: NULL;	CYCLIN-DEPENDENT KINASE 6; CHAIN: A; P19INK4D; CHAIN: B;	CYCLIN-DEPENDENT KINASE 6; CHAIN: A; PI9INK4D; CHAIN: B;	CYCLIN-DEPENDENT KINASE 6 INHIBITOR; CHAIN: A:	CYCLIN-DEPENDENT
SeqFold score			56.93						
PMF		-		-	-	-	-	0.3	96.0
Verify score		0.71		0.43	0.21	0.44	0.44	0.24	0.25
PSI- BLAST		2.60E-27	3.60E-29	7.20E-27	3.60E-29	2.60E-26	5.40E-29	3.60E-29	5.40E-34
End		212	201	182	160	202	148	291	162
Start AA		22	38	43	6	43	6	. 107	9
Chain D		en en en en en en en en en en en en en e				ø	В	. ✓	A
PDB ID		lawc	8pq1	1P48	1bd8	1bix	1blx	1bu9	1bu9
SEQ Sign	·	223	723	723	723	723	723	723	723

PDB annotation	INK4C; CELL CYCLE INHIBITOR, P18INK4C, TUMOR, SUPPRESSOR, CYCLIN- 2 DEPENDENT KINASE, HORMONE/GROWTH FACTOR	HORMONE/GROWTH FACTOR P18- INK4C; CELL CYCLE INHIBITOR, P18INK4C; TUMOR, SUPPRESSOR, CYCLIN-2 DEPENDENT KINASE, HORMONE/GROWTH FACTOR	CELL CYCLE INHIBITOR P18- INK4C(INK6), CELL CYCLE INHIBITOR, P18-INK4C(INK6), ANKYRIN REPEAT, 2 CDK 4/6 INHIBITOR	CELL CYCLE INHIBITOR P18- INK4C(INK6); CELL CYCLE INHIBITOR, P18-INK4C(INK6), ANKYRIN REPEAT, 2 CDK 4/6 INHIBITOR	TRANSCRIPTION FACTOR P65, P50D; TRANSCRIPTION FACTOR, IKBNFKB COMPLEX	TRANSCRIPTION FACTOR P65; P50D; TRANSCRIPTION FACTOR, IKBNFKB COMPLEX	TRANSCRIPTION FACTOR P65; P50D; TRANSCRIPTION FACTOR, IKBNFKB COMPLEX	TRANSCRIPTION FACTOR P65; P50D; TRANSCRIPTION FACTOR, IKB/NFKB COMPLEX	COMPLEX (TRANSCRIPTION REG/ANK REPEAT) COMPLEX (TRANSCRIPTION REGULATION/ANK REPEAT), ANKYRIN 2 REPEAT HELIX	COMPLEX (TRANSCRIPTION REGIANK REPEAT) COMPLEX
Coumpound	KINASE 6 INHIBITOR; CHAIN: A;	CYCLIN-DEPENDENT KINASIE 6 INHIBITOR; CHAIN: A;	CYCLIN-DEPENDENT KINASE 6 INHIBITOR; CHAIN: A, B;	CYCLIN-DEPENDENT KINASE 6 INHIBITOR; CHAIN: A, B;	NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF-KAPPA-B P50D SUBUNIT; CHAIN: C; I- KAPPA-B-ALPHA; CHAIN: D;	NF-KAPPA-B P65 SUBUNIT: CHAIN: A; NF-KAPPA-B P50D SUBUNIT; CHAIN: C; I- KAPPA-B-ALPHA; CHAIN: D;	NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF-KAPPA-B P50D SUBUNIT; CHAIN: C; I- KAPPA-B-ALPHA: CHAIN: D:	NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NP-KAPPA-B P50D SUBUNIT; CHAIN: C; I- KAPPA-B-ALPHA: CHAIN: D:	NF-KAPPA-B P65; CHAIN: A, C; NF-KAPPA-B P50; CHAIN: B, D; I-KAPPA-B-ALPHA; CHAIN: E, F:	NF-KAPPA-B P65; CHAIN: A, C; NF-KAPPA-B P50; CHAIN:
SeqFold score						10.19				
PMF score		0.88	0.35	0.99	-		0.95	0.43	0.65	_
Verify score		0.32	0.07	0.15	0.13		0.37	-0.02	0.06	0.31
PSI- BLAST		1.80E-26	1.80E-28	1.80E-33	1.80E-43	1.80E-43	3.60E-36	1.10E-28	5.40E-30	9.00E-44
End		234	284	191	174	208	229	280	288	174
Start		73	107	9	2	2	35	89	136	7.
Chain		∢	∢	∢	Q	Q	Ω	Ω	ш	மு
PDB ID		1 bu9	lihb	1ihb	likn	likn	1 ikn	lika	1nfi	1nfi
SEQ Seq		723	723	723	723	723	723	723	723	723

PDB annotation	(TRANSCRIPTION REGULATION/ANK REPEAT), ANKYRIN 2 REPEAT HELIX	COMPLEX (TRANSCRIPTION REG/ANK REPEAT) COMPLEX (TRANSCRIPTION REGULATION/ANK REPEAT), ANKYRIN 2 REPEAT HELIX	COMPLEX (TRANSCRIPTION REG/ANK REPEAT) COMPLEX (TRANSCRIPTION REGULATION/ANK REPEAT), ANKYRIN 2 REPEAT HELIX	COMPLEX (INHIBITOR/NUCLEASE) COMPLEX (INHIBITOR/NUCLEASE), COMPLEX (RI-ANG), HYDROLASE 2 MOLECULAR RECOGNITION, EPITOPE MAPPING, LEUCINE-RICH 3 REPEATS	COMPLEX (INHIBITOR/NUCLEASE) COMPLEX (INHIBITOR/NUCLEASE), COMPLEX (RI-ANG), HYDROLASE 2 MOLECULAR RECOGNITION, EPITOPE MAPPING, LEUCINE-RICH 3 REPEATS	COMPLEX (INHIBITOR/NUCLEASE) COMPLEX (INHIBITOR/NUCLEASE), COMPLEX (RI-ANG), HYDROLASE 2 MOLECULAR RECOGNITION, EPITOPE MAPPING, LEUCINE-RICH 3 REPEATS	COMPLEX (INHIBITOR/NUCLEASE) COMPLEX (INHIBITOR/NUCLEASE), COMPLEX (RI-ANG), HYDROLASE 2 MOLECULAR RECOGNITION, EPITOPE MAPPING, LEUCINE-RICH 3 REPEATS	COMPLEX (INHIBITORNUCLEASE) COMPLEX (INHIBITORNUCLEASE), COMPLEX (RI-ANG), HYDROLASE 2 MOLECULAR RECOGNITION, EPITOPE MAPPING, LEUCINE-RICH 3 REPEATS
Coumpound	B, D; I-KAPPA-B-ALPHA; CHAIN: E, F;	NF-KAPPA-B P65; CHAIN: A, C; NF-KAPPA-B P50; CHAIN: B, D; I-KAPPA-B-ALPHA; CHAIN: E, F;	NF-KAPPA-B P65; CHAIN: A, C; NF-KAPPA-B P50; CHAIN: B, D; I-KAPPA-B-ALPHA; CHAIN: E, F;	RIBONUCLEASE INHIBITOR; CHAIN: A, D, ANGIOGENIN; CHAIN: B, E;	RIBONUCLEASE INHIBITOR; CHAIN: A. D. ANGIOGENIN; CHAIN: B, E;	RIBONUCLEASE INHIBITOR; CHAIN: A, D, ANGIOGENIN; CHAIN: B, E;	RIBONUCLEASE INHIBITOR; CHAIN: A, D; ANGIOGENIN; CHAIN: B, E;	RIBONUCLEASE INHIBITOR; CHAIN: A, D; ANGIOGENIN; CHAIN: B, E;
SeqFold score			59.74				93.46	·
PMF		-		0.07	9.16	0.55	-	_
Verify score		0.49		-0.4	-0.3	-0.33		0.15
PSI- BLAST		5.40E-36	9.00E-44	1.30E-13	9.00E-17	2.60E-22	3.90E-36	9.10E-39
End AA		229	246	369	577	265	558	376
Start AA		34	36	-	219	267	82	83
Chain D		Э	ធ	·	∢	Ą	A	¥
PDB ID		Infi	lnfi	1a4y	1a4y	la4y	1a4y	la4y
SEQ ID NO:		723	723	725	725	725	725	725

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PDB annotation	COMPLEX (NUCLEAR PROTEIN/RNA) COMPLEX (NUCLEAR PROTEIN/RNA), RNA, SNRNP,RIBONUCLEOPROTEIN	COMPLEX (NUCLEAR PROTEIN/RNA) COMPLEX (NUCLEAR PROTEIN/RNA), RNA, SNRNP,RIBONUCLEOPROTEIN	COMPLEX (NUCLEAR PROTEIN/RNA) COMPLEX (NUCLEAR PROTEIN/RNA), RNA, SNRNP,RIBONUCLEOPROTEIN	COMPLEX (NUCLEAR PROTEIN/RNA) COMPLEX (NUCLEAR PROTEIN/RNA), RNA, SNRNP, RIBONUCLEOPROTEIN	COMPLEX (NUCLEAR PROTEIN/RNA) COMPLEX (NUCLEAR PROTEIN/RNA), RNA, SNRNP, RIBONUCLEOPROTEIN	COMPLEX (NUCLEAR PROTEIN/RNA) COMPLEX (NUCLEAR PROTEIN/RNA), RNA, SNRNP,RIBONUCLEOPROTEIN	COMPLEX (NUCLEAR PROTEIN/RNA) COMPLEX (NUCLEAR PROTEIN/RNA), RNA, SNRNP.RIBONUCLEOPROTEIN	COMPLEX (NUCLEAR PROTEIN/RNA) COMPLEX (NUCLEAR PROTEIN/RNA), RNA, SNRNP, RIBONICLEOPROTEIN	COMPLEX (NUCLEAR PROTEINRNA) COMPLEX (NUCLEAR PROTEINRNA), RNA, SNRNP, RIBONUCLEOPROTEIN	COMPLEX (MUCLEAR PROTEIN/RNA) COMPLEX (MUCLEAR PROTEIN/RNA), RNA, SNRNP, RIBONICLEOPROTEIN	COMPLEX (NUCLEAR PROTEIN/RNA) COMPLEX (NUCLEAR PROTEIN/RNA), RNA. SNRNP RIBONICLEOPROTEIN	CELL ADHESION LEUCINE RICH REPEAT, CALCIUM BINDING, CELL ADHESION	CELL ADHESION LEUCINE RICH REPEAT, CALCIUM BINDING, CELL ADHESION	CELL ADHESION LEUCINE RICH REPEAT, CALCIUM BINDING, CELL
Coumpound	U2 RNA HAIRPIN IV; CHAIN: Q, R; U2 A'; CHAIN: A, C; U2 B"; CHAIN: B, D;	U2 RNA HAIRPIN IV; CHAIN: Q, R; U2 A'; CHAIN: A, C; U2 B"; CHAIN: B, D;	U2 RNA HAIRPIN IV; CHAIN: Q, R; U2 A; CHAIN: A, C; U2 B"; CHAIN: B, D;	U2 RNA HAIRPIN IV; CHAIN: Q, R; U2 A; CHAIN: A, C; U2 B"; CHAIN: B, D;	U2 RNA HAIRPIN IV; CHAIN: Q, R; U2 A'; CHAIN: A, C; U2 B"; CHAIN: B, D;	U2 RNA HAIRPIN IV; CHAIN: Q, R; U2 A'; CHAIN: A, C; U2 B"; CHAIN: B, D;	U2 RNA HAIRPIN IV; CHAIN: Q, R; U2 A; CHAIN: A, C; U2 B"; CHAIN: B, D;	U2 RNA HAIRPIN IV; CHAIN: Q, R; U2 A; CHAIN: A, C; U2 B": CHAIN: B. D:	U2 RNA HAIRPIN IV; CHAIN: Q, R; U2 A; CHAIN: A, C; U2 B": CHAIN: B. D:	UZ RNA HAIRPIN IV; CHAIN: Q, R; U2 A'; CHAIN: A, C; U2 B": CHAIN: B. D:	U2 RNA HAIRPIN IV; CHAIN: Q, R; U2 A; CHAIN: A, C; U2 B": CHAIN: B. D:	INTERNALIN B; CHAIN: A;	INTERNALIN B; CHAIN: A;	INTERNALIN B; CHAIN: A;
SeqFold score														
PMF	0.57	0.75	0.77	0.45	0.99	0.84	0.88	69.0	0.45	0.55	96:0	. 89.0	1	1
Verify score	-0.02	90.00	0.12	-0.33	0.22	-0.06	0.24	-0.04	0	0.05	0.27	-0.09	0.02	0.12
PSI- BLAST	5.20E-27	1.30E-25	3.90E-19	6.50E-20	3.90E-29	1.20E-27	2.60E-25	3.90E-19	6.50E-20	9.10E-26	6.50E-29	1.60E-26	1.30E-24	5.40E-29
End	306	351	576	576	228	319	351	576	576	213	236	158	319	364
Start	134	203	430	454	68	134	203	430	454	82	68	_	104	216
Chain D	4	¥	∢	4	¥	ပ	ပ ·	၁	U	၁	ပ	Ą	¥	4
PDB UD	1a9n	1a9n	1a9n	1a9n	1a9n	1a9n	1a9n	189n	la9n	1a9n	la9n	140b	1406	1406
SEQ NO.	725	725	725	725	725	725	725	725	725	725	725	725	725	725

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PDB annotation	ADHESION	CELL ADHESION LEUCINE RICH REPEAT, CALCIUM BINDING, CELL ADHESION	CELL ADHESION LEUCINE RICH REPEAT, CALCIUM BINDING, CELL ADHESION	CELL ADHESION LEUCINE RICH REPEAT, CALCIUM BINDING, CELL ADHESION	TRANSFERASE CRYSTAL STRUCTURE, RAB GEBANYI GEBANYI TRANSFERASE	2.0 A 2 RESOLUTION, N-	FORMYLMETHIONINE, ALPHA SUBUNIT, BETA SUBUNIT		TRANSFERASE CRYSTAL	STRUCTURE, RAB	OEKAN I LUEKAN I LI KANSFEKASE,	FORMYLMETHIONINE. ALPHA	SUBUNIT, BETA SUBUNIT	TIMOSTAC TO LATITUTE THE	TRANSFERASE CRYSTAL STRUCTHIRE RAR	GERANYLGERANYLTRANSFERASE.	2.0 A 2 RESOLUTION, N-	FORMYLMETHIONINE, ALPHA	SUBUNIT, BETA SUBUNIT	TRANSFERASE CRYSTAL	STRUCTURE, RAB	GERANYLGERANYLTRANSFERASE,	2.0 A 2 RESOLUTION, N-	FORMYLMETHIONINE, ALPHA	SUBUNIT, BETA SUBUNIT		TRANSFERASE CRYSTAL	STRUCTURE, RAB GERANYLGERANYLTRANSFERASE,
Coumpound		INTERNALIN B; CHAIN: A;	INTERNALIN B; CHAIN: A;	INTERNALIN B; CHAIN: A;	RAB GERANYLGERANYLTRANSF PRASE AI PHA STIRITMIT:	CHAIN: A, C; RAB	GERANYLGERANYLTRANSF ERASE BETA SUBUNIT:	CHAIN: B, D;	RAB	GERANYLGERANYLTRANSF	CHAIN A C. RAR	GERANYLGERANYLTRANSF	ERASE BETA SUBUNIT;	CHAIN: B, D;	RAB GERANYI GERANYI TRANSF	ERASE ALPHA SUBUNIT:	CHAIN: A, C; RAB	GERANYLGERANYLTRANSF	ERASE BETA SUBUNIT;	RAB	GERANYLGERANYLTRANSF	ERASE ALPHA SUBUNIT;	CHAIN: A, C; RAB	GERANYLGERANYLTRANSF	ERASE BETA SUBUNIT;	CHAIN: B, D;	RAB	GERANYLGERANYLTRANSF ERASE ALPHA SUBUNIT;
SeqFold score																												
PMF		0.4	0.92	0.34	I				0.95					,00	0.96					0.22							0.07	
Verify score		0.04	0.18	0.09	0.12				-0.23					9	87.0					0.08							0.07	
PSI- BLAST		9.00E-28	5.40E-24	1.10E-21	3.60E-13				5.40E-13					21 200 0	9.00E-13					1.80E-13							7.20E-09	
End AA		406	166	969	301				346					9,0	707					533						į	100	
Start		242	32	415	216				246					970	807					428							4/2	
Chain		∢	⋖	e e	Ä				Ą						<					A							⋖_	
ED ED		1406	140b	140b	1dce				1dce					1	98					Idee							Jace	
SEQ NO:		725	725	. 725	725				725					13.6	3					725		_				, ,	3	

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PDB annotation	2.0 A 2 RESOLUTION, N- FORMYLMETHIONINE, ALPHA SUBUNIT, BETA SUBUNIT	TRANSFERASE CRYSTAL STRUCTURE, RAB GERANYLGERANYLTRANSFERASE, 2.0 A 2 RESCLUTION, N- FORMYLMETHIONINE, ALPHA SUBUNIT, BETA SUBUNIT	CONTRACTILE PROTEIN LEUCINE- RICH REPEAT, BETA-BETA-ALPHA CYLINDER, DYNEIN, 2 CHLAMYDOMONAS, FLAGELLA	CONTRACTILE PROTEIN LEUCINE- RICH REPEAT, BETA-BETA-ALPHA CYLINDER, DYNEIN, 2 CHLAMYDOMONAS, FLAGELLA	CONTRACTILE PROTEIN LEUCINE- RICH REPEAT, BETA-BETA-ALPHA CYLINDER, DYNEIN, 2 CHLAMYDOMONAS, FLAGELLA	CONTRACTILE PROTEIN LEUCINE- RICH REPEAT, BETA-BETA-ALPHA CYLINDER, DYNEIN, 2 CHLAMYDOMONAS, FLAGEILA	CONTRACTILE PROTEIN LEUCINE- RICH REPEAT, BETA-BETA-ALPHA CYLINDER, DYNEIN, 2 CHLAMYDOMONAS, FLAGELLA	LIGASE CYCLIN A/CDK2- ASSOCIATED PROTEIN P45; CYCLIN A/CDK2-ASSOCIATED PROTEIN P19; SKP1, SKP2, F-BOX, LRN, LEUCINE- RICH REPEAT, SCF, UBIQUITIN, 2 E3, UBIQUITIN PROTEIN LIGASE	LIGASE CYCLIN A/CDK2- ASSOCIATED PROTEIN P45; CYCLIN A/CDK2-ASSOCIATED PROTEIN P19; SKP1, SKP2, F-BOX, LRR, LEUCINE-
Coumpound	CHAIN: A, C; RAB GERANYLGERANYLIRANSF ERASE BETA SUBUNIT; CHAIN: B, D;	RAB GERANYLGERANYLTRANSF ERASE ALPHA SUBUNIT; CHAIN: A, C; RAB GERANYLGERANYLTRANSF ERASE BETA SUBUNIT; CHAIN: B, D;	OUTER ARM DYNEIN; CHAIN: A;	SKP2; CHAIN: A, C, E, G, I, K, M, O; SKP1; CHAIN: B, D, F, H, J, L, N, P;	SKP?; CHAIN: A, C, E, G, I, K, M, O; SKPI; CHAIN: B, D, F, H, J, L, N, P;				
SeqFold		l.							-
PMF score	-	0.95	0.21	0.94	0.12	0.52	0.75	0.04	60:00
Verify score		0.3	-0.34	-0.36	-0.73	-0.47	-0.79	0.02	-0.35
PSI- BLAST		1.30E-13	3.60E-13	1.10E-14	7.20E-13	1.30E-13	3.90E-26	3.60E-07	1.30E-09.
End AA		170	140	324	346	164	213	172	575
Start		09	17	210	232	70	82	-	264
Chain			¥	¥	ď	٧	4	∢	A
PDB ID		1 dce	1ds9	1ds9	1ds9	1ds9	· 1ds9	lfqv	lfqv
SEQ S D S		725	725	725	725	725	725	725	725

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PDB annotation	RICH RÉPEAT, SCF, UBIQUITIN, 2 E3, UBIQUITIN PROTEIN LIGASE	LIGASE CYCLIN A/CDK2- ASSOCIATED PROTEIN P45; CYCLIN ACDK2-ASSOCIATED PROTEIN P19; SKP1, SKP2, F-BOX, LRR, LEUGINE- RICH REPEAT; SCF, UBIQUITIN, 2 E3, UBIQUITIN PROTEIN LIGASE	LIGASE CYCLIN A/CDK2- ASSOCIATED P45; CYCLIN A/CDK2- ASSOCIATED P19; SKP1, SKP2, F-BOX, LRN3, LEUCINE-RICH REPEATS, SCF, 2 UBIQUITIN, E3, UBIQUITIN PROTEIN LIGASE	LIGASE CYCLIN A/CDK2- ASSOCIATED P45; CYCLIN A/CDK2- ASSOCIATED P19; SKP1, SKP2, F-BOX, LRRS, LEUCINE-RICH REPEATS, SCF, 2 UBIQUITIN, E3, UBIQUITIN PROTEIN LIGASE	LIGASE CYCLIN A/CDK2- ASSOCIATED P45: CYCLIN A/CDK2- ASSOCIATED P19; SKP1, SKP2, F-BOX, LRRS, LEUCINE-RICH REPEATS, SCF, 2 UBIQUITIN, E3, UBIQUITIN PROTEIN LIGASE	TRANSCRIPTION RNAIP; RANGAP; GTPASE-ACTIVATING PROTEIN FOR SPII, GTPASE-ACTIVATING PROTEIN, GAP, RNAIP, RANGAP, LRR, LEUCINE- 2 RICH REPEAT PROTEIN, TWINNING, HEMIHEDRAL TWINNING, 3 MEROHEDRAL TWINNING, MEROHEDRAL TWINNING, MEROHEDRAL	ACETYLATION RNASE INHIBITOR, RIBONUCLEASE/ANGIOGENIN INHIBITOR ACETYLATION, LEUCINE- RICH REPEATS	ACETYLATION RNASE INHIBITOR, RIBONUCLEASE/ANGIOGENIN INHIBITOR ACETYLATION LEUCINE-
Coumpound		SKP2; CHAIN: A, C, B, G, I, K, M, O; SKP1; CHAIN: B, D, F, H, J, L, N, P;	SKP?; CHAIN: A, C; SKP1; CHAIN: B, D;	SKP2; CHAIN: A, C; SKP1; CHAIN: B, D;	SKP2; CHAIN: A, C; SKP1; CHAIN: B, D;	GTPASE-ACTIVATING PROTEIN RNA1_SCHPO; CHAIN: A, B;	RIBONUCLEASE INHIBITOR; CHAIN: NULL;	RIBONUCLEASE INHIBITOR; CHAIN: NULL;
SeqFold score		·						
PMF score		0.06	0.11	0.27	0.31	0.04	0.22	0.87
Verify score		-0.06	-0.09	-0.12	-0.14	-0.22	-0.07	-0.26
PSI- BLAST		1.00E-16	3.60E-11	7.80E-19	6.50E-40	1.30E-09	1.10E-19	5.40E-21
End		253	453	577	338	431	498	592
Start AA		84	244	401	68	211	110	239
Chain TO		∢	∢	∀ .	∢ .	∢		
PDB CD		1fqv	16.2	162	1fs2	lyrg	2bnh	2bnh
SEQ ID NO:		725	725	725	725	725	725	725

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PDB annotation	RICH REPEATS	ACETYLATION RNASE INHIBITOR, RIBONUCLEASE/ANGIOGENIN INHIBITOR ACETYLATION, LEUCINE- RICH REPEATS	ACETYLATION RNASE INHIBITOR, RIBONUCLEASE/ANGIOGENIN INHIBITOR ACETYLATION, LEUCINE- RICH REPEATS	1 Common and the second second	COMPLEX (TRANSCRIFTION REGULATION/DNA) GABPALPHA; GABPBETAI; COMPLEX	(TRANSCRIPTION	KEGULATION/DNA), DNA-BINDING, Z NUCLEAR PROTEIN, ETS DOMAIN,	ANK YKIN KEFEAIS, IKANSCKIFIION 3 FACTOR	COMPLEX (TRANSCRIPTION REGULATION/DNA) GABPALPHA;	GABPBETA1; COMPLEX	(TRANSCRIPTION	REGULATION/DNA), DNA-BINDING, 2	NUCLEAR PROTEIN, ETS DOMAIN,	ANKYKIN KEPEATS, TKANSCKIPTION 3 FACTOR	COMPLEX (TRANSCRIPTION	REGULATION/DNA) GABPALPHA;	GABFBETAI; COMPLEX	REGULATION/DNA), DNA-BINDING, 2	NUCLEAR PROTEIN, ETS DOMAIN,	ANKYRIN REPEATS, TRANSCRIPTION	3 FACTOR	COMPLEX (TRANSCRIPTION PEGIT ATTOMONA) CARPAI PHA:	GABPBETA1: COMPLEX	TRANSCRIPTION	REGULATION/DNA), DNA-BINDING, 2	NUCLEAR PROTEIN, ETS DOMAIN, ANKYRIN REPEATS, TRANSCRIPTION
Coumpound		RIBONUCLEASE INHIBITOR; CHAIN: NULL;	RIBONUCLEASE INHIBITOR; CHAIN: NULL;		GA BINDING PROTEIN ALPHA; CHAIN: A; GA BINDING PROTEIN BETA 1;	CHAIN: B; DNA; CHAIN: D, E;			GA BINDING PROTEIN ALPHA: CHAIN: A: GA	BINDING PROTEIN BETA 1;	CHAIN: B; DNA; CHAIN: D, E;				GA BINDING PROTEIN	ALPHA; CHAIN: A; GA	BINDING PROTEIN BELA I;	בתיקווי. B, בויס, כווסווי. ב, ב,				GA BINDING PROTEIN	RINDING PROTEIN BETA 1.	CHAIN: B; DNA; CHAIN: D, E;		•
SeqFold			90.38												53.5						١					
PMF score		0.01			-0.03				0.29									-		_		0.72				
Verify score		-0.33			0.12				0.19				_					_				0.32				
PSI- BLAST		1.80E-19	1.10E-19		3.60E-32				1.30E-35						1.30E-35							1.60E-31				
End		448	558		145				.167						179							203				
Start AA		35	88		7				23						27				ś			55				
Chain ID			,		ф				В						В							Д				•
PDB		2bnh	2bnh		lawc				lawc						lawc							lawc				
SEQ D	ä	725	725		728				728						728							728		_		

PDB annotation	3 FACTOR	HORMONE/GROWTH FACTOR P18- INK4C; CELL CYCLE INHIBITOR, P18INK4C; TUMOR, SUPPRESSOR, CYCLIN- 2 DEPENDENT KINASE, HORMONE/GROWTH FACTOR	HORMONE/GROWTH FACTOR P18- INK4C; CELL CYCLE INHIBITOR, P18INK4C, TUMOR, SUPPRESSOR, CYCLIN- 2 DEPENDENT KINASE, HORMONE/GROWTH FACTOR	HORMONE/CROWTH FACTOR P18- INK4C; CELL CYCLE INHIBITOR, P18INK4C; TUMOR, SUPPRESSOR, CYCLIN- 2 DEPENDENT KINASE, HORMONE/GROWTH FACTOR	CELL CYCLE INHIBITOR P18- INK4C(INK6); CELL CYCLE INHIBITOR, P18-INK4C(INK6), ANKYRIN REPEAT, 2 CDK 4/6 INHIBITOR	TRANSCRIPTION FACTOR P65; P50D; TRANSCRIPTION FACTOR, IKBNFKB COMPLEX	TRANSCRIPTION FACTOR P65; P50D; TRANSCRIPTION FACTOR, IKBNFKB COMPLEX	COMPLEX (TRANSCRIPTION REG/ANK REPEAT) COMPLEX (TRANSCRIPTION REGULATION/ANK REPEAT), ANKYRIN 2 REPEAT HELIX	COMPLEX (TRANSCRIPTION REG/ANK REPEAT) COMPLEX (TRANSCRIPTION REGULATION/ANK REPEAT), ANKYRIN 2 REPEAT HELIX	COMPLEX (TRANSCRIPTION REGIANK REPEAT) COMPLEX (TRANSCRIPTION REGULATION/ANK REPEAT), ANKYRIN 2 REPEAT HELIX
Coumpound		CYCLIN-DEPENDENT KINASE 6 INHIBITOR; CHAIN: A;	CYCLIN-DEPENDENT KINASE 6 INHIBITOR; CHAIN: A;	CYCLIN-DEPENDENT KINASE 6 INHIBITOR; CHAIN: A;	CYCLIN-DEPENDENT KINASE 6 INHIBITOR; CHAIN: A, B;	NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF-KAPPA-B P50D SUBUNIT; CHAIN: C; I- KAPPA-B-ALPHA; CHAIN: D;	NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF-KAPPA-B P50D SUBUNIT; CHAIN: C; I- KAPPA-B-ALPHA; CHAIN: D;	NF-KAPPA-B P65; CHAIN: A, C; NF-KAPPA-B P50; CHAIN: B, D; I-KAPPA-B-ALPHA; CHAIN: E, F;	NF-KAPPA-B P65; CHAIN: A, C; NF-KAPPA-B P50; CHAIN: B, D; I-KAPPA-B-ALPHA; CHAIN: E, F;	NF-KAPPA-B P65; CHAIN: A, C; NF-KAPPA-B P50; CHAIN: B, D; I-KAPPA-B-ALPHA; CHAIN: E, F;
SeqFold score								ř	&	
PMF		0.01	0.09	-0.15	0.41	0.24	0.23	0.22	0.36	60.0
Verify score		0.33	0.28	0.04	0.25	0.15	0.27	0.24	0.16	0.03
PSI- BLAST		1.80E-29	1.805-29	1.40E-27	3.60E-29	9.00E-37	1.80E-32	9.00E-37	5.40E-32	1.30E-26
End		183	150	250	166	162	198	162	198	245
Start		23	5	97	23	12	18	12	17	50
Chain		<	V	<	∢	Ω	Q	ш	ω	<u>ы</u> .
PDB		1bu9	1bu9	1bu9	lihb	1 ikm	1 ikn	Inf	Infi	Infi
SEQ D	ä	728	728	728	728	728	728	728	728	728

PDB annotation	COMPLEX (TRANSCRIPTION REGIANK REPEAT) COMPLEX (TRANSCRIPTION REGULATION/ANK REPEAT), ANKYRIN 2 REPEAT HELIX	PHOSPHOTRIESTERASE PHOSPHOTRIESTERASE, HYPOTHETICAL PROTEIN	SI RNA-BINDING DOMAIN POLYRIBONUCLEOTIDE NUCLEOTIDYLTRANSFERASE, SI RNA-BINDING DOMAIN, POLYNUCLEOTIDE PHOSPHORYLASE 2 (PNPASE)	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX
Coumpound	NF-KAPPA-B P65; CHAIN: A, C; NF-KAPPA-B P50; CHAIN: B, D; I-KAPPA-B-ALPHA; CHAIN: E, F;	PHOSPHOTRIESTERASE HOMOLOGY PROTEIN; CHAIN: A, B;	PNPASE; CHAIN: NULL;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN: CHAIN: C, F, G;
SeqFold score					81.11				
PMF	0.03	69.0	-	0.82		-	6.0	0.96	1
Verify score	0.06	0.37	0.13	0.08		-0.04	-0.33	0.06	0.27
PSI- BLAST	5.40E-29	3.90E-07	5.40E-23	1.10E-26	7.80E-45	7.80E-45	6.50E-41	1.80E-43	3.60E-46
End	253	208	618	217	443	469	498	217	245
Start AA	88	74	540	137	361	389	417	136	164
Chain	跙	4		¥.	¥	٧	٧	ပ	C
PDB UD	11g	1bf6	1sro	lalh	lalh	lalh	laih	Jme y	1me y
SEQ S B S	728	729	731	736	736	736	736 .	736	736

PDB annotation	(ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA FINGER, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA FINGER, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
Coumpound		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C. F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A. B, D, E; CONSENSUS ZINC FINGER PROTEIN: CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;
SeqFold score							104.61		! !
PMF		-	-	_	-	ı		-	
Verify score		0.41	0.19	0.38	0.26	0.46		0.44	0.05
PSI- BLAST		9.00E-48	7.20E-49	9.00E-51	7.20E-51	3.60E-50	7.20E-51	9.00E-50	3.60E-50
End		273	301	357	385	413	414	144	469
. Start AA		192	220	276	304	332	332	360	388
Chain		U	U	U	ပ	U	U	U	U
PDB	1	Jme y	1me y	J mc	Jme y	1me y	1me y	lme y	y y
SEQ	ÿ	736	736	736	736	736	736	736	736

					· · · · · · · · · · · · · · · · · · ·			
PDB annotation		COMPLEX (ZINC FINGERDNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGERDNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX TRANSCRIPTION REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATIONDNA) COMPLEX (TRANSCRIPTION REGULATIONDNA), RNA POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATIONDNA) COMPLEX (TRANSCRIPTION REGULATIONDNA), RNA POLYMERASE III, 2 TRANSCRIPTION INTIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATIONDNA) COMPLEX (TRANSCRIPTION REGULATIONDNA), RNA POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATIONDNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1. ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX
Coumpound		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	TFIIIA; CHAIN: A, D: 5S RIBOSOMAL RNA GENE; CHAIN: B, C. E, F;	TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	TFIIIA, CHAIN: A, D, 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA: CHAIN: A, B;
SeqFold	score				112.4			
PMF	score	-	0.33	-		0.95	0.92	0.19
Verify	score	-0.1	0.09	0.04		-0.11	-0.04	0.01
PSI-	BLAST	1.80E-50	1.40E-34	1.10E-36	2.60E-79	3.60E-38	1.60E-35	5.40E-29
End	\$	497	282	338	444	479	499	217
Start	ΨΨ	416	137	193	276	333	361	E
Chain	В	U	4	4	A	V V	4	O
aud	a a	y y	1466	146	1466	931	146	lubd
Cas	B	736 736	736	736	736	736	736	736

PDB annotation	(TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2	FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION.	INITIATOR ELEMENT, YYI, ZINC 2 FINGER PROTEIN, DNA-PROTEIN	RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1;	TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2	FINGER PROTEIN, DNA-PROTEIN	RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA) YING-YANG 1;	INITIATOR ELEMENT, YY1, ZINC 2	FINGER PROTEIN, DNA-PROTEIN	(TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1;	TRANSCRIPTION INITIATION,	INITIATOR ELEMENT, YY1, ZINC 2	FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX	(TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA) YING-YANG I;	INITIATOR ELEMENT, YYI, ZINC 2 FINGER PROTEIN, DNA-PROTEIN
Coumpound		YYI; CHAIN. C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;		YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 BUILT ATOR ET EMENT DNA:	CHAIN: A, B;		YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS P5	NITIATOR ELEMENT DNA;			YY1; CHAIN: C; ADENO-	ASSOCIATED VIRUS P5 INITIATOR PI EMENT DNA:	CHAIN: A, B;	•		YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS P5	INITIATOR ELEMENT DNA;	CHAIN: A, B;			YY1; CHAIN: C; ADENO-	ASSOCIATED VIRUS PS	CHAIN: A, B;
SeqFold score										,						89.47	•						
PMF		-		_			_				-										_		
Verify score		0.03		0.18			0.11				0.1						,				10.0		
PSI- BLAST		1.00E-56		1.30E-55			1.20E-57				3.90E-57					3.90E-57					2.60E-56		
End		273		329			358		_		385					414					469		
Start		162		218			246				274					304					359		
Chain		O		O			U				O					ပ					U		
PDB	1	1ubd		lubd			lubd				1ubd					lubd					Jubd		
SEQ	ë Z	736		736			736		_		736					736		_			736		

PDB annotation	RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DAY YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATIONDNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INTITATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA).	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA).	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLJ; GLJ, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI,
Coumpound		YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C,
SeqFold score									
PMF score		, 	96.0	66:0	-	1	-	-	-
Verify score		0	-0.18	-0.04	0.2	0.38	0.22	0.03	0.58
PSI- BLAST		1.80E-34	3.90E-51	3.60E-34	2.60E-58 ·	1.80E-34	2.60E-70	5.20E-74	1.30E-73
End		469	497	497	275	300	303	359	387
Start		368	386	396	150	164	164	192	249
Chain		O	O	o ·	∀	4	∢	∢	4
PDB TD		lubd	lubd	lubd	2gli	2gli	2gli	2gli	2gli
SEQ B	ğ	736	736	736	736	736	736	736	736

	NA-	er gli; gli, dna-	ER GLI; GLI, ONA-	ER GLI; GLI, DNA-	ER GLI; GLI, ONA-		ER GLI; GLI, DNA-	ER GLI; GLI, DNA- ER GLI; GLI, DNA-	ER GLI; GLI, DNA- ER GLI; GLI, DNA-	ER GLI; GLI, DNA- ER GLI; GLI, DNA- ATALYTIC PTIDE) STIDE) STIDE EENT,	ER GLI; GLI, DNA- ER GLI; GLI, DNA- ATALYTIC ATALYTIC FITDE) IENT, EX O, BETA	ER GLI; GLI, DNA- ER GLI; GLI, DNA- ATALYTIC ATALYTIC FIENT, EENT, SX NT
PDB annotation	ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEINONA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)		COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLJ; GLJ, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA) COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GL. ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA) COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GL. ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA) CATALYTIC ANTIBODY CATALYTIC ANTIBODY, ESTERASE	COMPLEX (DNA-BINDING PROTEINDINA) FIVE-FINGER GLI ZINC FINGER, COMPLEX (DNA- BINDING PROTEINIDING COMPLEX (DNA-BINDING PROTEINIDING PROTEINIDING PROTEINIDING ZINC FINGER, COMPLEX (DNA- BINDING PROTEINIDINA) CATALYTIC ANTIBODY CATALY ANTIBODY, ESTERASE COMPLEX (ANTIBODY/PEPTIDE) POLYSPECIFICITY, CROSS REACTIVITY, FAB-FRAGMENT, PEPTIDE, 2 HIV-1, COMPLEX (ANTIBODY/PEPTIDE)	COMPLEX (DNA-BINDING PROTEINDNA) FIVE-FINGER GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEINDNA) BINDING PROTEINDNA) PROTEINDNA) FIVE-FINGER GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEINDNA) CATALYTIC ANTIBODY CATALY CATALYTIC ANTIBODY (PEPTIDE) POLYSPECIFICITY, CROSS REACTIVITY, FAB-FRAGMENT, PEPTIDE, 2 HIV-1, COMPLEX (ANTIBODY/PEPTIDE) CONTRACTILE PROTEIN GONTRACTILE PROTEIN BARNEL BARNEL	COMPLEX (DNA-BINDING PROTEINDING) PROTEINDING) PROFELS (DNA-BINDING) PROTEINDING) PROTEINDING) PROTEINDING) PROTEINDING) PROTEINDING) PROTEINDING) PROTEINDING) PROTEINDING) PROTEINDING) PROTEINDING) PROTEINDING) PROTEINDING) PROTEINDING) PROTEINDING) PROTEINDING) PROTEINDING) PROTEINDING) PROTEINDING) PROTEING) PROTEING) PROTEING) PROTEING) PROTEING) PROTEING) PROTEING) PROTEING) BARREL ANTIBODY FAB FRAGMENT ANTIBODY FAB FRAGMENTATIBODY FAB F
Coumpound		IC FINGER PROTEIN GLI1; AIN: A; DNA; CHAIN: C,	IC FINGER PROTEIN GLII; AIN: A; DNA; CHAIN: C,	AC FINGER PROTEIN GLJ I; AIN: A: DNA; CHAIN: C,	NC FINGER PROTEIN GLII; IAIN: A; DNA; CHAIN: C,	-	AC FINGER PROTEIN GLII; AIN: A; DNA; CHAIN: C,	AC FINGER PROTEIN GLII; AIN: A; DNA; CHAIN: C, AC FINGER PROTEIN GLII; AIN: A; DNA; CHAIN: C,	= =	ii ii j	F F 7	F F F
SeqFold score	ά	92.84 ZIN CH CH CH	ZZ CH D; CH	Z G	E CE		35°C	1	25.38 S5.38			
PMF S	-	6			· · ·			0.03				
Verify score	-		-0.06	0.06	0.03		0.19					
PSI- BLAST		5.20E-74	2.60E-72	3.60E-33	2.60E-68	1	3.60E-34					
End		387	471	468	499	1	496	216	216	230	216 230 230 212	230 230
Start		250	305	340	360		368	898	368	368	368 68 68 19 19 126	368
Chain		<	4	4	4		4	4 4	4 4 7	4 4 1 4	4 4 4	4 4 1 4 4 1
PDB ID		2gli	2gli	2gli	2gli		2gli	2gli 2gli	2gli 2gli 1a0q	2gli 2gli 1a0q 1b0g	2gli 2gli 1a0q 1bog 1fhg	2gli 2gli 1a0q 1b0g 1fhg
SEQ	Ö	736	736	736	736		736	736	736	736	736 741 741 741	736 741 741 741 741

PDB annotation					COMPLEX (IMMUNOGLOBULIN/LIPOPROTEIN) OSPA; COMPLEX (IMMUNOGLOBULIN/LIPOPROTEIN), OUTER SURFACE 2 PROTEIN A COMPLEXED WITH FAB184.1, BORRELIA BURGDORFERI 3 STRAIN B31			CELL ADHESION PROTEIN NCAM MODULE 2; CELL ADHESION, GLYCOPROTEIN, HEPARIN-BINDING, GPI-ANCHOR, 2 NEURAL ADHESION MOLECULE, IMMUNOGLOBULIN FOLD, HOMOPHILIC 3 BINDING, CELL ADHESION PROTEIN	LIGASE CYCLIN A/CDK2- ASSOCIATED PROTEIN P45; CYCLIN A/CDK2-ASSOCIATED PROTEIN P19; SKP1, SKP2, F-BOX, LRR, LEUCINE- RICH REPEAT, SCF, UBIQUITIN, 2 E3, UBIQUITIN PROTEIN LIGASE
Coumpound	GLYCOPROTEIN CD2 (HUMAN) IHNF 3	T LYMPHOCYTE ADHESION GLYCOPROTEIN CD2 (FUMAN) 1HNF 3	T LYMPHOCYTE ADHESION GLYCOPROTEIN CD2 (RAT) 1HNG 3	T LYMPHOCYTE ADHESION GLYCOPROTEIN CD2 (RAT) 1HNG 3	FAB 184.1; CHAIN: L, H; OUTER SURFACE PROTEIN A; CHAIN: O;	MUSCLE PROTEIN TITIN MODULE M5 (CONNECTIN) ITMM 3 (LMR, MNIMIZED AVERAGE STRUCTURE) ITNM 4 ITNM 58	IMMUNOGLOBULIN IGG2A FAB FRAGMENT (CNI206) 2GFB 3	NEURAL CELL ADHESION MOLECULE, LARGE ISOFORM; CHAIN: A;	SKP2; CHAIN: A, C, E, G, I, K, M, O; SKP1; CHAIN: B, D, F, H, I, L, N, P;
SeqFold score			69.09		55.92		53.4		
PMF		0.76		0.35		-0.18		-0.18	0.92
Verify score		60.0		0.35	·	0.09		0	-0.07
PSI- BLAST		9.10E-18	2.60E-18	2.60E-18	0.0011	3.60E-14	3.60E-05	1.80E-09	2.60E-05
End		199	217	210	230	213	230	84	81
Start AA		28	25	52	61	131	61	52	46
Chain ID			Ą	V	'n		V	∢	¥
PDB		1hnf	1hng.	[hng	losp	Tom -	2gfb	3nc m	Ifqv
SEQ NO.	2	741	741	741	741	741	741	741	747

	,				———			 1	
PDB annotation	COMPLEX (DNA-BINDING PROTEIN/DNA) UPSTREAM STIMULATORY FACTOR 1: USF, DNA BINDING, BASIC-HELIX-LOOP-HELIX, LEUCINE ZIPPER, 2 TRANSCRIPTION FACTOR, COMPLEX (DNA-BINDING) PROTEIN/DNA)	TRANSCRIPTION FACTOR P65; P50D; TRANSCRIPTION FACTOR, IKBINFKB COMPLEX	COMPLEX (TRANSCRIPTION FACTOR/DNA) NF-KB P50, COMPLEX (TRANSCRIPTION FACTOR/DNA)		HYDROLASE MALTOGENIC ALPHA AMYLASE, AMYLASE, GLYCOSIDE HYDROLASE, STARCH DEGRADATION	-	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZNC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX
Coumpound	USF; CHAIN: A, B; DNA; CHAIN: C, D;	NF.KAPPA-B P65 SUBUNIT; CHAIN: A; NP-KAPPA-B P50D SUBUNIT; CHAIN: C; I- KAPPA-B-ALPHA; CHAIN: D;	NUCLEAR FACTOR KAPPA- B; CHAIN: A, B; KB SITE, DNA (5'-D(TGAGAATTCCC)- 3'); CHAIN: C, D;	GLYCOSYLTRANSFERASE CYCLODEXTRIN GLUCANOTRANSFERASE (E.C.2.4.1.19) (CGTASE) 1CYG	APHA-AMYLASE; CHAIN: A;	CHROMOSOMAL PROTEIN UBIQUITIN 1UBI 3	QGSR ZINC FINGER PEPTIDE: CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE: CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;
SeqFold score				·					
PMF	0.09	0.41	0.04	0.05	0.37	0.03	0.03	0.03	0.04
Verify	-0.61	0.39	-0.37	-0.02	0.11	-0.58	-0.45	-0.33	-0.52
PSI- BLAST	0.0078	0.0013	0.0013	0.0016	0.0061	0.0065	3.60E-25	3.60E-24	1.405-44
End	390	346	319	1059	196	1617	314	162	220
Start	353	243	234	932	854	1529	229	78	134
Chain TD	V.	<	V		4		4	V	U
PDB ID	lan4	1ik	lnfk	lcyg	1qho	lubi	falh	laih	Ime y
SEQ NO:	750	750	750	754	754	754	756	756	756

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PDB annotation	(ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CXYSTAL STRUCTURE, COMPLEX	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION. PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CXYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CEYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CXYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
Coumpound		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;
SeqFold score			·						
PMF score		90.0	0.28	0.12	-0.07	0.17	0.05	0.07	-0.19
Verify score		-0.6	-0.15	-0.19	0.06	-0.44	0.06	-0.3	0.06
PSI- BLAST		9.00E-44	3.60E-42	1.10E-42	3.60E-30	1.80E-42	5.40E-12	9.00E-11	9.00E-11
End		.284	465	612	627	162	190	284	
Start		193	379	526	555	77	163	254	437
Chain		ပ	O	ပ	U	ပ	5	ົນ	g
PDB ID		y y	Jme y	Jme y	y y	lme y	Jme y	Ime y	Jme y
SEQ D		756	756	756	756	756	756	952	756

PDB annotation	COMPLEX (TRANSCRIPTION REGULATION/DNA) TFIIIA; 5S GENE; NMR, TFIIIA, PROTEIN, DNA, TRANSCRIPTION FACTOR, 5S RNA 2 GENE, DNA BINDING PROTEIN, ZINC FINGER, COMPLEX 3 (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INITATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN. DNA-PROTEIN RECOGNITION, 3 COMPLEX TRANSCRIPTION, REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATIONDNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX CTRANSCRIPTION REGILIATIONNA)	COMPLEX (TRANSCRIPTION REGULATIONDNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX. THE ANGEIPTION, 3 COMPLEX.	COMPLEX (TRANSCRIPTION
Conmpound	TRANSCRIPTION FACTOR IIIA; CHAIN: A; 5S RNA GENE; CHAIN: E, F;	TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI, CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YY1; CHAIN: C; ADENO-
SeqFold score		69.84				,	
PMF	0.34		0.11	0.11	0.23	0	0.01
Verify score	-0.43		-0.18	-0.77	-0.21	-0.72	-0.53
PSI- BLAST	1.80E-15	1.40E-34	1.40E-34	1.305-30	1.80E-27	1.80E-31	1.80E-30
End AA	432	285	302	253	436	578	130
Start AA	349	106	135	139	325		85
Chain D	∢	<	Ą	ပ	ပ	υ .	O
PDB ID	9	115	1116	PgnI	Iubd	pqn	Inbd
SEQ NO D	756	756	756	756		756	756

				,	 						,					
PDB annotation	REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	TRANSCRIPTION REGULATION TRANSCRIPTION REGULATION, ADRI, ZINC FINGER, NMR	TRANSCRIPTION REGULATION TRANSCRIPTION REGULATION, ADRI, ZINC FINGER, NMR	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	STRUCTURAL PROTEIN TWO REPEATS OF SECTRIN AT PHA	HELICAL LINKER REGION, 22	TANDEM 3-HELIX COILED-COILS, STRUCTURAL PROTEIN	STRUCTURAL PROTEIN TWO	REPEATS OF SPECTRIN, ALPHA HELICAL LINKER REGION, 2.2 TANDEM 3-HELIX COILED-COILS, STRUCTURAL PROTEIN	CYTOSKELETON	SIGNALING PROTEIN DAPPI, PHISH, BAM32: PLECKSTRIN, 3-	PHOSPHOINOSITIDES, INOSITOL	TELKAKISPHUSPHATE Z SIGNAL TRANSDUCTION PROTEIN, ADAPTOR PROTEIN	SIGNALING PROTEIN ARFI GUANINE	NUCLEOTIDE EXCHANGE FACTOR AND PH DOMAIN	TRANSFERASE RECEPTOR TYROSINE KINASE, PROTEIN INTERACTION
Coumpound	ASSOCIATED VIRUS P5 INITATOR ELEMENT DNA; CHAIN: A, B;	ADRI; CHAIN: NULL;	ADRI; CHAIN: NULL;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ALPHA SPECTRIN; CHAIN:	S G	•	ALPHA SPECTRIN; CHAIN:	,4, B, C;	BETA-SPECTRIN; IDRO 6 CHAIN: NULL; IDRO 7	DUAL ADAPTOR OF PHOSPHOTYROSINE AND 3-	CHAIN: A;		GRP1; CHAIN: A;		EPHA4 RECEPTOR TYROSINE KINASE; CHAIN:
SeqFold score	·															
PMF		0.11	90:04	0.07	-0.02			0		0.43	68'0			89.0		-0.09
Verify score		-0.63	-0.13	-0.53	0.12			0.13		0.08	0.44	,		0.56		0.04
PSI- BLAST		3.60E-15	1.40E-15	1.10E-28	7.80E-14			9.10E-15		5.20E-06	5.20E-18			1.20E-18		3.60E-12
End AA		406	554	286	726			749		1203	1203			1203		857
Start AA		349	498	142	537			995		1096	1093			1097		
Chain ID				¥	<			V			∢			A		∢
PDB ID		2adr	2adr	2gli	lcm			1cun		1dro	1 1 68			-Ifgy		X O O
SEQ NO:		756	756	756	757			757		757	757			757		762

	MODULE, 2 DIMERIZATION DOMAIN, TRANSFERASE		ISBUCTION SAM I RECEPTOR, SIGNAL ON, OLIGOMER	SIGNAL TRANSDUCTION SAM DOMAIN, EPH RECEPTOR, SIGNAL TRANSDUCTION, OLIGOMER TYROSINE-PROTEIN KINASE NMR, RECEPTOR OLIGOMERIZATION, EPH RECEPTORS, TYROSINE 2 PHOSPHORYLATION, SIGNAL TRANSDUCTION, TYROSINE-PROTEIN 3 KINASE	ISDUCTION SAM I RECEPTOR, SIGNAL ON, OLIGOMER COTEN KINASE NMR, LIGOMERIZATION, EPH TYROSINE 2 ATION, SIGNAL ION, TYROSINE-PROTEIN	ISDUCTION SAM I RECEPTOR, SIGNAL ON, OLIGOMER GOTEIN KINASE NMR, LIGOMERIZATION, EPH TYROSINE 2 ATION, SIGNAL ON, TYROSINE-PROTEIN OSPHOLIPID BINDING	SIGNAL TRANSDUCTION SAM DOMAIN, EPH RECEPTOR, SIGNAL TRANSDUCTION, OLIGOMER TYROSINE-PROTEIN KINASE NMR, RECEPTOR OLIGOMERIZATION, EPH RECEPTOR, TYROSINE 2 PHOSPHORYLATION, SIGNAL TRANSDUCTION, TYROSINE-PROTEIN 3 KINASE CALCIUM/PHOSPHOLIPID BINDING PROTEIN PII, CALPACTIN LIGHT CHAIN; S100 FAMILY, EF-HAND PROTEIN, LIGAND OF ANNEXIN II, 2	ISDUCTION SAM I RECEPTOR, SIGNAL ON, OLIGOMER COTEIN KINASE NMR, LIGOMERIZATION, EPH TYROSINE 2 ATION, SIGNAL ON, TYROSINE-PROTEIN OSPHOLIPID BINDING CALPACTIN LIGHT FAMILY, EF-HAND SAND OF ANNEXIN II, 2 OSPHOLIPID BINDING	ISDUCTION SAM IRECEPTOR, SIGNAL ON, OLIGOMER. GOTEIN KINASE NAR. LIGOMERIZATION, EPH TYROSINE 2 ATION, SIGNAL ON, TYROSINE-PROTEIN OSPHOLIPID BINDING CALPACTIN LIGHT FAMILY, EF-HAND SAND OF ANNEXIN II, 2 OSPHOLIPID BINDING CALPACTIN LIGHT CALPACTIN LIGHT CALPACTIN LIGHT OSPHOLIPID BINDING CALPACTIN LIGHT CALPACTIN LIGHT	ISDUCTION SAM IRECEPTOR, SIGNAL ON, OLIGOMER GOTEIN KINASE NMR, LIGOMERIZATION, EPH TYROSINE 2 ATION, SIGNAL ON, TYROSINE-PROTEIN CALPACTIN LIGHT FAMILY, EF-HAND SAND OF ANNEXIN II, 2 OSPHOLIPID BINDING CALPACTIN LIGHT FAMILY, FF-HAND AND OF ANNEXIN II, 2 OSPHOLIPID BINDING OSPHOLIPID BINDING CALPACTIN LIGHT FAMILY, EF-HAND FAMILY, EF-HAND	SIGNAL TRANSDUCTION SAM DOMAIN, EPH RECEPTOR, SIGNAL TRANSDUCTION, OLIGOMER TYROSINE-PROTEIN KINASE NMR, RECEPTOR, OLIGOMERLZATION, EPH RECEPTOR, TYROSINE 2 PHOSPHORYLATION, SIGNAL TRANSDUCTION, TYROSINE-PROTEIN 3 KINASE CALCIUM/PHOSPHOLIPID BINDING PROTEIN, LIGAND OF ANNEXIN II, 2 CALCIUM/PHOSPHOLIPID BINDING PROTEIN, LIGAND OF ANNEXIN II, 2 CALCIUM/PHOSPHOLIPID BINDING PROTEIN CHAIN; SIOO FAMILY, EF-HAND PROTEIN CHAIN; SIOO FAMILY, EF-HAND PROTEIN PII, CALPACTIN LIGHT CHAIN; SIOO FAMILY, EF-HAND PROTEIN, LIGAND OF ANNEXIN II, 2 PROTEIN, LIGAND OF ANNEXIN II, 2	SIGNAL TRANSDUCTION SAM DOMAIN, EPH RECEPTOR, SIGNAL TRANSDUCTION, OLIGOMER TYROSINE-PROTEIN KINASE NMR, RECEPTOR, OLIGOMERIZATION, EPH RECEPTOR, TYROSINE 2 PHOSPHORYLATION, SIGNAL TRANSDUCTION, TYROSINE-PROTEIN 3 KINASE CALCIUM/PHOSPHOLIPID BINDING PROTEIN LIGAND OF ANNEXIN II, 2 CALCIUM/PHOSPHOLIPID BINDING PROTEIN CALCIUM/PHOSPHOLIPID BINDING PROTEIN CALCIUM/PHOSPHOLIPID BINDING CALCIUM/PHOSPHOLIPID BINDING CALCIUM/PHOSPHOLIPID BINDING CALCIUM/PHOSPHOLIPID BINDING CALCIUM/PHOSPHOLIPID BINDING PROTEIN LIGAND OF ANNEXIN II, 2 CALCIUM/PHOSPHOLIPID BINDING PROTEIN, SIOU FAMILY, EF-HAND PROTEIN LIGAND OF ANNEXIN II, 2 CALCIUM/PHOSPHOLIPID BINDING PROTEIN LIGAND OF ANNEXIN II, 2 CALCIUM/PHOSPHOLIPID BINDING PROTEIN	ISDUCTION SAM IRECEPTOR, SIGNAL ON, OLIGOMER ACTEN KINASE NAR, LIGOMERIZATION, EPH TYROSINE 2 ATION, SIGNAL ON, TYROSINE-PROTEIN OSPHOLIPID BINDING CALPACTIN LIGHT FAMILY, EF-HAND SAND OF ANNEXIN II, 2 OSPHOLIPID BINDING CALPACTIN LIGHT FAMILY, EF-HAND SAND OF ANNEXIN II, 2 OSPHOLIPID BINDING CALPACTIN LIGHT FAMILY, EF-HAND SAND OF ANNEXIN II, 2 OSPHOLIPID BINDING CALPACTIN LIGHT FAMILY, EF-HAND SAND OF ANNEXIN II, 2 OSPHOLIPID BINDING COSPHOLIPID BINDING COSPHOLIPID BINDING	ISDUCTION SAM IRECEPTOR, SIGNAL ON, OLIGOMER OTHORIZATION, EPH TYROSINE 2 ATION, SIGNAL ON, TYROSINE-PROTEIN OSPHOLIPID BINDING CALPACTIN LIGHT FAMILY, EF-HAND SAND OF ANNEXIN 11, 2 OSPHOLIPID BINDING CALPACTIN LIGHT FAMILY, EF-HAND SAND OF ANNEXIN 11, 2 OSPHOLIPID BINDING CALPACTIN LIGHT FAMILY, EF-HAND SAND OF ANNEXIN 11, 2 OSPHOLIPID BINDING CALPACTIN LIGHT FAMILY, EF-HAND SAND OF ANNEXIN 11, 2 OSPHOLIPID BINDING CALPACTIN LIGHT FAMILY, EF-HAND SAND OF ANNEXIN 11, 2 OSPHOLIPID BINDING NORMALY TRICE.	SIGNAL TRANSDUCTION SAM DOMAIN, EPH RECEPTOR, SIGNAL TRANSDUCTION, OLIGOMER TRANSDUCTION, OLIGOMER RECEPTOR OLIGOMERIZATION, EPH RECEPTOR OLIGOMERIZATION, EPH RECEPTOR, TYROSINE 2 PHOSPHORYLATION, SIGNAL TRANSDUCTION, TYROSINE-PROTEIN SKINASE CALCIUMPHOSPHOLIPID BINDING PROTEIN LIGAND OF ANNEXIN II, 2 CALCIUMPHOSPHOLIPID BINDING PROTEIN CALCIUMPHOSPHOLIPID BINDING PROTEIN CALCIUMPHOSPHOLIPID BINDING PROTEIN CALCIUMPHOSPHOLIPID BINDING PROTEIN CALCIUMPHOSPHOLIPID BINDING PROTEIN CALCIUMPHOSPHOLIPID BINDING PROTEIN CALCIUMPHOSPHOLIPID BINDING PROTEIN CALCIUMPHOSPHOLIPID BINDING CALCIUMPHOSPHOLIPID BINDING CALCIUMPHOSPHOLIPID BINDING CALCIUMPHOSPHOLIPID BINDING CALCIUMPHOSPHOLIPID BINDING CALCIUMPHOSPHOLIPID BINDING CALCIUMPHOSPHOLIPID BINDING CALCIUM-BINDING PROTEIN CALCIUM-BINDING PROTEIN CALCIUM-BINDING PROTEIN CALCIUM-BINDING PROTEIN CALCIUM-BINDING PROTEIN CALCIUM-BINDING PROTEIN CALCIUM-BINDING CERTUM TRIC-	ISDUCTION SAM IRECEPTOR, SIGNAL ON, OLIGOMER COTEIN KINASE NAR, LIGOMERIZATION, EPH TYROSINE 2 ATION, SIGNAL ON, TYROSINE-PROTEIN OSPHOLIPID BINDING CALPACTIN LIGHT FAMILY, EF-HAND JAND OF ANNEXIN II, 2 OSPHOLIPID BINDING CALPACTIN LIGHT FAMILY, EF-HAND SAND OF ANNEXIN II, 2 OSPHOLIPID BINDING CALPACTIN LIGHT FAMILY, EF-HAND SAND OF ANNEXIN II, 2 OSPHOLIPID BINDING NORMAN TRIC- SIDURS I - 75; CERLUM- LIUM-BINDING	ISDUCTION SAM IRECEPTOR, SIGNAL ON, OLIGOMER ACTEIN KINASE NAR, LIGOMERIZATION, EPH TYROSINE 2 ATION, SIGNAL ON, TYROSINE-PROTEIN OSPHOLIPID BINDING CALPACTIN LIGHT FAMILY, EF-HAND AND OF ANNEXIN II, 2 OSPHOLIPID BINDING CALPACTIN LIGHT FAMILY, EF-HAND SAND OF ANNEXIN II, 2 OSPHOLIPID BINDING CALPACTIN LIGHT FAMILY, EF-HAND OSPHOLIPID BINDING CALPACTIN LIGHT FAMILY, EF-HAND SAND OF ANNEXIN II, 2 OSPHOLIPID BINDING NOERUW TRIC- SIDUES I - 75; CERLUM- LCIUM-BINDING LCIUM-BINDING LCIUM-BINDING	ISDUCTION SAM IRECEPTOR, SIGNAL ON, OLIGOMER OTEN KINASE NAR, LIGOMERIZATION, EPH TYROSINE 2 ATION, SIGNAL ON, TYROSINE-PROTEIN OSPHOLIPID BINDING CALPACTIN LIGHT FAMILY, EF-HAND IAND OF ANNEXIN II, 2 OSPHOLIPID BINDING CALPACTIN LIGHT FAMILY, EF-HAND SAND OF ANNEXIN II, 2 OSPHOLIPID BINDING CALPACTIN LIGHT FAMILY, EF-HAND SAND OF ANNEXIN II, 2 OSPHOLIPID BINDING CALPACTIN LIGHT FAMILY, EF-HAND SAND OF ANNEXIN II, 2 OSPHOLIPID BINDING CALPACTIN LIGHT FAMILY, FF-HAND SAND OF ANNEXIN II, 2 COSPHOLIPID BINDING COSPHOLIPID BINDING ING PROTEIN LCIUM-BINDING LCIUM-BINDING ING PROTEIN S100B, ING PROTEIN S100B, ING PROTEIN S100B, ING PROTEIN S100B, ING PROTEIN S100B, ING PROTEIN S100B, INGRETA, S100B, NARR,	SIGNAL TRANSDUCTION SAM DOMAIN, EPH RECEPTOR, SIGNAL TRANSDUCTION, OLIGOMER TYROSINE-PROTEIN KINASE NAR, RECEPTOR OLIGOMERIZATION, EPH RECEPTOR, TYROSINE 2 PHOSPHORYLATION, SIGNAL TRANSDUCTION, TYROSINE-PROTEIN SKINASE CALCIUM/PHOSPHOLIPID BINDING PROTEIN, LIGAND OF ANNEXIN II, 2 CALCIUM/PHOSPHOLIPID BINDING PROTEIN CALCIUM/PHOSPHOLIPID BINDING PROTEIN CALCIUM/PHOSPHOLIPID BINDING PROTEIN CALCIUM/PHOSPHOLIPID BINDING PROTEIN CALCIUM/PHOSPHOLIPID BINDING PROTEIN CALCIUM/PHOSPHOLIPID BINDING PROTEIN CALCIUM/PHOSPHOLIPID BINDING PROTEIN CALCIUM/PHOSPHOLIPID BINDING PROTEIN CALCIUM/PHOSPHOLIPID BINDING PROTEIN CALCIUM/PHOSPHOLIPID BINDING PROTEIN CALCIUM/PHOSPHOLIPID BINDING PROTEIN CALCIUM/PHOSPHOLIPID BINDING PROTEIN LOADED, CALCIUM-BINDING PROTEIN METAL BINDING PROTEIN S100B, S100BETA, S100BETA, S100B, NMR, S100BETA, S100BETA, S100B, NMR, S100BETA, S100BETA, S100B, NMR,	ISDUCTION SAM IRECEPTOR, SIGNAL ON, OLIGOMER, JGOMERIZATION, EPH TYROSINE 2 ATION, SIGNAL ON, TYROSINE-PROTEIN OSPHOLIPID BINDING CALPACTIN LIGHT FAMILY, EF-HAND OSPHOLIPID BINDING CALPACTIN LIGHT FAMILY, EF-HAND OSPHOLIPID BINDING OSPHOLIPID BINDING OSPHOLIPID BINDING CALPACTIN LIGHT FAMILY, EF-HAND AND OF ANNEXIN II, 2 OSPHOLIPID BINDING CALPACTIN LIGHT CALLACTIN LIGHT CALLACTIN LIGHT CALLACTIN LIGHT CALLACTIN LIGHT CALLACTIN LIGHT CALLACTIN LIGHT CALLACTIN LIGHT CALLACTIN LIGHT CALLACTIN LIGHT CALLACTIN LIGHT CALLACTIN LIGHT CALLACTIN LIGHT CALLACTIN SIOOB, ING PROTEIN ING PROTEIN ING PROTEIN ING PROTEIN SIOOB,	ISDUCTION SAM IRECEPTOR, SIGNAL ON, OLIGOMER, JGOMERIZATION, EPH TYROSINE 2 ATION, SIGNAL ON, TYROSINE-PROTEIN OSPHOLIPID BINDING CALPACTIN LIGHT FAMILY, EF-HAND OSPHOLIPID BINDING OSPHOLIPID BINDING CALPACTIN LIGHT FAMILY, EF-HAND OSPHOLIPID BINDING OSPHOLIPID BINDING CALPACTIN LIGHT FAMILY, EF-HAND AND OF ANNEXIN II, 2 OSPHOLIPID BINDING CALPACTIN LIGHT FAMILY, EF-HAND AND OF ANNEXIN II, 2 OSPHOLIPID BINDING CALCIUM-BINDING ING PROTEIN NOTERUM TRIC- SIDURG 1-75; CERLUM- LCIUM-BINDING ING PROTEIN SIOOB, ING PROTEIN SIOOB, UNG-HELLINGS, EF-HAND, SIOO ALCIUM-BINDING UNC-HELLINGS, EF-HAND, SIOO ALCIUM-BINDING UNC-HELLINGS, EF-HAND, SIOO ALCIUM-BINDING	ISDUCTION SAM IRECEPTOR, SIGNAL ON, OLIGOMER, COTEIN KINASE NAR, LIGOMERIZATION, EPH TYROSINE 2 ATION, SIGNAL ON, TYROSINE-PROTEIN OSPHOLIPID BINDING CALPACTIN LIGHT FAMILY, EF-HAND OSPHOLIPID BINDING OSPHOLIPID BINDING OSPHOLIPID BINDING CALPACTIN LIGHT FAMILY, EF-HAND OSPHOLIPID BINDING OSPHOLIPID BINDING CALPACTIN LIGHT FAMILY, EF-HAND COSPHOLIPID BINDING CALPACTIN LIGHT CALPACTIN LIGHT CALPACTIN LIGHT CALLACTIN LIGHT CALLACTIN LIGHT CALLACTIN LIGHT CALLACTIN LIGHT CALLACTIN LIGHT CALLACTIN LIGHT CALLACTIN LIGHT CALLACTIN LIGHT CALLACTIN LIGHT CALLACTIN SIOOB, UNCHELIX BUNDIG, MENSIONAL CALLIN BUNDICE, MENSIONAL	SIGNAL TRANSDUCTION SAM JONARIN, EPH RECEPTOR, SIGNAL TRANSDUCTION, OLIGOMER, TRANSDUCTION, OLIGOMER, TREOSINE-PROTEIN KINASE NAR, ECEPTOR, CLIGOMERZATION, EPH REANSDUCTION, TYROSINE 2 HOSPHORYLATION, SIGNAL RANSDUCTION, TYROSINE 2 HOSPHORYLATION, SIGNAL RANSDUCTION, TYROSINE 2 HOSPHORYLATION, SIGNAL SALCIUMPHOSPHOLIPID BINDING PROTEIN, LIGAND OF ANNEXIN II, 2 CALCIUM/PHOSPHOLIPID BINDING PROTEIN CALCIUM/PHOSPHOLIPID BINDING PROTEIN CALCIUM/PHOSPHOLIPID BINDING PROTEIN CALCIUM/PHOSPHOLIPID BINDING PROTEIN CALCIUM/PHOSPHOLIPID BINDING PROTEIN CALCIUM/PHOSPHOLIPID BINDING PROTEIN CALCIUM/PHOSPHOLIPID BINDING PROTEIN CALCIUM/PHOSPHOLIPID BINDING PROTEIN CALCIUM/PHOSPHOLIPID BINDING PROTEIN METAL BINDING PROTEIN SIOOB, SIOOBETA, SIOOBETA, SIOOB, NAR, DEPOLAR COUPLINGS, EF-HAND, SIOO PROTEIN, FOUR-HELIX BINDING PROTEIN, FOUR BINDING PROTEIN, FOUR-HELIX BINDING PROTEIN FOUR-HELIX BINDING	ISDUCTION SAM IRECEPTOR, SIGNAL ON, OLIGOMER, IGOMERIZATION, EPH TYROSINE 2 ATION, SIGNAL ON, TYROSINE-PROTEIN OSPHOLIPID BINDING CALPACTIN LIGHT FAMILY, EF-HAND AND OF ANNEXIN II, 2 OSPHOLIPID BINDING CALPACTIN LIGHT FAMILY, EF-HAND OSPHOLIPID BINDING CALPACTIN LIGHT FAMILY, EF-HAND OSPHOLIPID BINDING CALPACTIN LIGHT FAMILY, EF-HAND OSPHOLIPID BINDING CALCIUM-BINDING NO CERUM TRIC- SIDUES 1 - 75; 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GOTEIN KINASE NAR. LIGOMERIZATION, EPH TYROSINE 2 ATION, SIGNAL ON, TYROSINE-PROTEIN OSPHOLIPID BINDING CALPACTIN LIGHT FAMILY, EF-HAND SAND OF ANNEXIN II, 2 OSPHOLIPID BINDING CALPACTIN LIGHT FAMILY, EF-HAND SAND OF ANNEXIN II, 2 OSPHOLIPID BINDING CALPACTIN LIGHT FAMILY, EF-HAND AND OF ANNEXIN II, 2 OSPHOLIPID BINDING CALPACTIN LIGHT FAMILY, EF-HAND AND OF ANNEXIN II, 2 OSPHOLIPID BINDING LCIUM-BINDING ING PROTEIN S100B, UNCHELIX BINDING ALCIUM-BINDING ON-HELIX BINDING ALCIUM-BINDING ALCIUM-BINDING ALCIUM-BINDING ON-HELIX BUNDLE, GENSIONAL SOLUTION STRUCTURE
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A; EPHB2; CHAIN: A, B, C, D, E, F, G, H;	PHB2; CHAIN: A, B, G, H;		EPHRIN TYPE-B RECEPTOR 2; CHAIN: NULL;			S100A10; CHAIN: A, B;	100A10; CHAIN: A, E	100A10; CHAIN: A, E	\$100A10; CHAIN: A, B;	100A10; CHAIN: A, E	100A10; CHAIN: A, E	100A10; CHAIN: A, E	100A10; CHAIN: A, E 100A10; CHAIN: A, E ALMODULIN; CHAI	S100A10; CHAIN: A, B; S100A10; CHAIN: A, B; CALMODULIN; CHAIN: NULL;	100A10; CHAIN: A, E 100A10; CHAIN: A, E ALMODULIN; CHAI	100A10; CHAIN: A, E 100A10; CHAIN: A, E ALMODULIN; CHAI	S100A10; CHAIN: A, B; S100A10; CHAIN: A, B; CALMODULIN; CHAIN NULL; S-100 PROTEIN, BETA	100A10; CHAIN: A, E 100A10; CHAIN: A, E ALMODULIN; CHAI IULL; HAIN; CHAIN: A, B;	S100A10; CHAIN: A, B S100A10; CHAIN: A, B S100 PROTEIN, BETA CHAIN; CHAIN: A, B;	100A10; CHAIN: A, E 100A10; CHAIN: A, E ALMODULIN; CHAI IULL; HAIN; CHAIN: A, B;	100A10; CHAIN: A, E ALMODULIN; CHAI TULL; HAIN; CHAIN: A, BET,	100A10; CHAIN: A, E ALMODULIN; CHAI TULL; HAIN; CHAIN: A, BET	100A10; CHAIN: A, E 100A10; CHAIN: A, E ALMODULIN; CHA 100 PROTEIN, BET HAIN; CHAIN: A, B	S100A10; CHAIN: A, E S100A10; CHAIN: A, E CALMODULIN; CHAIN NULL; CHAIN; CHAIN: A, B CALCIUM-BINDING CALCIUM-BINDING	S100A10; CHAIN: A, B; S100A10; CHAIN: A, B; S100A10; CHAIN: A, B; NULL; CALMODULIN; CHAIN: NULL; CHAIN; CHAIN: A, B; CALCIUM-BINDING PROTEIN CALBINDIN D9K (INTACT FORM) (NMR. 13 STRLICTURES) 1, CB1 3
A; EPH	EPH	Б.	EPH 2; C		-												4	4	4	4	4	4	4	4	4
						69.57	69.57	69.57	69.57	69.57	69.57	69.57	69.57	69.57	69.57	69.57	69.57	69.57	69.57	145.44	145.44	145.44	145.44	145.44	145.44
		-0.02	-0.02									-	-011	-0.11	1 0-	0.11	-0.11	1.0-	0.11	-0.11	-0.11	-0.11	-0.11	-0.11	-0.11
		0.1	0.53						0.67	0.67	0.67	0.67	0.67	0.67	0.67	0.67	0.67	0.67	0.67	0.67	0.67	0.67	0.67	0.67	0.67
		1.10E-14	7.20E-14			1.30E-20	1.30E-20	1.30E-20	1.30E-20	1.30E-20	1.30E-20	1.30E-20	1.30E-20 1.30E-20 1.30E-20	1.30E-20 1.30E-20 3.60E-21	1.30E-20 1.30E-20	1.30E-20 1.30E-20 3.60E-21	1.30E-20 1.30E-20 3.60E-21	1.30E-20 1.30E-20 3.60E-21	1.30E-20 1.30E-20 3.60E-21	1.30E-20 1.30E-20 3.60E-21	1.30E-20 1.30E-20 3.60E-21	1.30E-20 1.30E-20 3.60E-21	1.30E-20 1.30E-20 3.60E-21	1.30E-20 1.30E-20 7.20E-19	1.30E-20 1.30E-20 3.60E-21 7.20E-19
		863	861		\top	32	+		+					+											
		793	797						2																
				_	1		+	1																	
_	-	1b4f A	1888	-	+	la4p A																		_ 	
	Ö	762	762	_	\dagger	763	++	++	++		- 	++	 	 	- 	 	++	 	- 		- 	- 	- - - - - - - - - - - - - - - - - - - 	- - - 	-

PDB annotation			CALCIUM-BINDING PROTEIN CALMODULIN APO TR2C-DOMAIN; ICMF 9	METAL TRANSPORT CALMODULIN, HIGH RESOLUTION, DISORDER	TRANSPORT PROTEIN CALCIUM BINDING, EF HAND, FOUR-HELIX BUNDLE	CALCIUM-BINDING CALCIUM- BINDING, ZINC, METAL-BINDING, ACETYLATION			CALCIUM-BINDING PROTEIN CALCIUM-BINDING PROTEIN, CALCIUM-DEPENDENT PROTEASE, APO 2 FORM, SMALL SUBUNIT	STRUCTURAL PROTEIN HELIX-TURN- HELIX	METAL TRANSPORT CALMODULIN, HIGH RESOLUTION, DISORDER	METAL BINDING PROTEIN YEAST FREOUENIN EF-HAND, CALCIUM	CALCIUM-REGULATED MUSCLE	CONTRACTION, MOSCLE CONTRACTION, CALCIUM-BINDING,	IROFONIN, E-F. HAND, 2 OPEN CONFORMATION REGULATORY DOMAIN, CALCIUM-REGULATED 3
Coumpound	PROTEIN CALBINDIN D9K (INTACT FORM) (NMR, 13 STRUCTURES) ICBI 3	CALCIUM-BINDING PROTEIN CALMODULIN COMPLEXED WITH CALMODULIN-BINDING DOMAIN OF ICDM 3 CALMODULIN-DEPENDENT PROTEIN KINASE II ICDM 4	CALMODULIN (VERTEBRATE); ICMF 6 CHAIN: NULL; ICMF 7	CALMODULIN; CHAIN: A;	CALMODULIN; CHAIN: A;	S-100 PROTEIN; CHAIN: NULL;	CONTRACTILE SYSTEM PROTEIN TROPONIN C 1TOP 3		CALPAIN; CHAIN: A, B;	CARDIAC TROPONIN C: CHAIN: A;	CALMODULIN; CHAIN: A;	CALCIUM-BINDING PROTEIN NCS-1; CHAIN: A:	TROPONIN C; CHAIN: NULL;		
SeqFold score						137.02									
PMF score		0.06	0.15	-0.17	0.18		-0.07		0.09	0.28	0.39	-0.03	0.47		
Verify score		0.21	0.17	90.0	-0.17		0.07		-0.18	-0.29	-0.18	0.05	-0.12		
PSI- BLAST		9.00E-21	7.20E-21	3.60E-23	7.20E-20	1.40E-16	7.20E-20		2.60E-09	2.60E-09	1.30E-09	2.60E-08	3.90E-10		
End		81	81	. 68	81	89	92		252	252	252	252	252		
Start AA		4	1	4	80	2	٧.		135	131	129	129	135		
Chain The Chain		∢		Ą	٧				∢	4	٧	¥			
PDB CI		E E	1cmf	lexr	1771	1mh o	Itop	1	Iajo	륟	lexr	1fpw	1tcf		
SEQ NO:		763	763	763	763	763	763		766	992	992	992	99/		

PDB annotation	MUSCLE CONTRACTION .	CALCIUM-BINDING PROTEIN EF. HAND 1TNX 14			MEMBRANE PROTEIN AQPI WATER CHANNEL, TWO-DIMENSIONAL CRYSTAL, ELECTRON 2 DIFFRACTION, ELECTRON MICROSCOPY	MEMBRANE PROTEIN AQPI WATER CHANNEL, TWO-DIMENSIONAL CRYSTAL, ELECTRON 2 DIFFRACTION, ELECTRON MICROSCOPY		ENDOCYTOSIS/EXOCYTOSIS SYNAPTOTAGMIN, CZ-DOMAIN, EXOCYTOSIS, NEUROTRANSMITTER Z RELEASE, ENDOCYTOSIS/EXOCYTOSIS	ENDOCYTOSIS/EXOCYTOSIS NSECI; PROTEIN-PROTEIN COMPLEX, MULTI-SUBUNIT	ENDOCYTOSIS/EXOCYTOSIS BETA SANDWICH, CALCIUM ION, C2 DOMAIN	TRANSFERASE CALCIUM++, PHOSPHOLIPID BINDING PROTEIN, CALCIUM-BINDING 2 PROTEIN, PHOSPHATIDYLSERINE, PROTEIN	
Coumpound		TROPONIN C; 1TNX 4 CHAIN: NULL; 1TNX 5	CALCIUM BINDING PROTEIN CALMODULIN (TR-22-C\$ FRAGMENT COMPRISING RESIDUES 78 - 148 ITRC 3 OF THE INTACT MOLECULE) 1TRC 4	CALCIUM-BINDING PROTEIN PARVALBUMIN (ALPHA LINEAGE) 5PAL 3	AQUAPORIN-1; CHAIN: A;	AQUAPORIN-1; CHAIN: A;		SYNAPTOTAGMIN I¸ CHAIN: A;	SYNTAXIN BINDING PROTEIN 1; CHAIN: A; SYNTAXIN 1A; CHAIN: B;	SYNAPTOTAGMIN III; CHAIN: A;	PROTEIN KINASE C, ALPHA TYPE; CHAIN: A;	CALCTUM/PHOSPHOLIPID
SeqFold score			-				•					
PMF		0.31	0.95	0.53	0.51	0.88		0.94	0.01	0.82	0.71	1
Verify score		-0.53	-0.37	0.27	-0.22	-0.13		0.21	-0.13	0.38	0.48	0.3
PSI- BLAST		1.00E-10	I.30E-08	1.30E-08	1.10E-73	3.90B-76		1.30E-11	5.20E-05	6.50E-07	0.0026	6.50E-12
End AA		252	252	252	227	227		664	252	647	999	664
Start AA		140	194	177		6		541	26	541	541	541
Chain D			∢		₹ .	∢		٧	В	A	∀	
PDB ID		1tmx	Itro	Spal	1fgy	1fqy		1 вуп	1dn1	1dqv	1dsy	Irsy
SEQ NO.		992	766	766		892		692	769	169	691	769

											
PDB annotation		P2I; SOS; COMPLEX (ONCOGENE PROTEIN/EXCHANGE PACTOR), SMALL GIPASE, 2 EXCHANGE FACTOR	P21; SOS; COMPLEX (ONCOGENE PROTEIN/EXCHANGE FACTOR), SMALL GTPASE, 2 EXCHANGE FACTOR	P21; SOS; COMPLEX (ONCOGENE PROTEIN/EXCHANGE PACTOR), SMALL GTPASE, 2 EXCHANGE FACTOR.	SIGNAL TRANSDUCTION PROTEIN	CYTOSKELETON		SIGNALING PROTEIN DAPPI, PHISH, BAM32; PLECKSTRIN, 3-PHOSPHOINOSITIDES, INOSITOL TETRAKISPHOSPHATE 2 SIGNAL TRANSDUCTION PROTEIN, ADAPTOR PROTEIN	SIGNALING PROTEIN DAPPI, PHISH, BAM32; PLECKSTRIN, 3-PHOSPHOINOSITIDES, INOSITOL TETRAKISPHOSPHATE 2 SIGNAL TRANSDUCTION PROTEIN, ADAPTOR PROTEIN	SIGNALING PROTEIN ARFI GUANINE NUCLEOTIDE EXCHANGE FACTOR AND PH DOMAIN	
Coumpound	BINDING PROTEIN SYNAPTOTAGMIN I (FIRST C2 DOMAIN) (CALB) IRSY 3	H-RAS, CHAIN: R; SON OF SEVENLESS-1; CHAIN: S;	H-RAS; CHAIN: R; SON OF SEVENLESS-1; CHAIN: S;	H-RAS, CHAIN: R; SON OF SEVENLESS-1; CHAIN: S;	BETA-SPECTRIN; 1BTN 4 CHAIN: NULL; 1BTN 5	BETA-SPECTRIN; 1DRO 6 CHAIN: NULL; 1DRO 7	SIGNAL TRANSDUCTION PROTEIN DYNAMIN (PLECKSTRIN HOMOLOGY DOMAIN) (DYNPH) 1DYN 3	DUAL ADAPTOR OF PHOSPHOTYROSINE AND 3- CHAIN: A;	DUAL ADAPTOR OF PHOSPHOTYROSINE AND 3- CHAIN: A;	GRP1; CHAIN: A;	PHOSPHORYLATION PLECKSTRIN (N-TERMINAL
SeqFold score		79.72									
PMF					99.0	0.4	0.25	6.9	0.05	0.36	0.24
Verify score		·	0.2	0.26	-0.09	-0.22	-0.04	0.3	0.04	0.35	-0.02
PSI- BLAST		9.10E-77	9.10E-77	7.20E-61	1.40E-18	7.20E-12	5.40E-06	1.00E-07	1.30E-12	1.10E-16	1.10E-13
End		311	291	312	541	546	541	546	539	546	542
Start		-	12	14	438	439	456	429	441	447	437
Chain ID		S	S	Ø			٧	∢	4	Ą	
EDB ID		1bkd	15kd	1bkd	1btn	1dro	Idyn	1168	1fb8	1fgy	1 pls
S B B S B S		770	770	170	770	770	770	770	770	770	770

						·		
PDB annotation	,	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 COYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CEYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC PINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CYYSTAL STRUCTURE, COMPLEX (ZINC PINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CYYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
Coumpound	PLECKSTRIN HOMOLOGY DOMAIN) MUTANT 1PLS 3 WITH LEU GLU (HIS)6 ADDED TO THE C TERMINUS 1PLS 4 (INS(G105-LEHHHHHH)) (NMR, 25 STRUCTURES) 1PLS 5	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE: CHAIN: B. C:	QGSR ZINC FINGER PEPTIDE, CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B. C;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, B; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN; A, B, D. E; CONSENSUS ZINC FINGER PROTEIN; CHAIN; C. F, Q;	DNA; CHAIN: A, B, D, E. CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;
SeqFold			78.23					
PMF		0.99			~	_	_	<u>-</u>
Verify score		0.19		0.37	0.29	0.46	0.29	0.25
PSI- BLAST		6.50E-41	5.20E-45	5.40E-46	3.60E-47	1.80E-48	1.10E-49	3.60E-51
End		210	351	209	237	265	293	377
Start AA		131	269	128	156	184	212	296
Chain		V	¥	υ .	ပ	Ų	ပ	ပ
PDB CD		lath	lalh	Ime y	lme y	Ime y	Jme y	lme y
SEQ NO EQ		772	27.2		772	27.2	772	27.

PDB annotation	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CEYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CXYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CYYSTAL STRUCTURE, COMPLEX ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CYYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (TRANSCRIPTION COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA REGULATION/DNA), RNA POLYMERAS B III, 2 TRANSCRIPTION NITTATION ATMOST FROM TRANSCRIPTION TRANSC	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA REGULATION/DNA), RNA PULYMERASE III, 2 TRANSCRIPTION PUTTA ATTON	INTERNATION, ZENC FINGER PROTEIN COMPLEX (TRANSCRIPTION (TRANSCRIPTION (TRANSCRIPTION REGULATION/DNA), RNA
Coumpound	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	TFIIIA; CHAM: A, D; 5S RIBOSOMAL RNA GENE; CHAM: B, C, E, F;	TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;
SeqFold score	102.88					115.72		
PMF		-		-	0.42		0.76	0.94
Verify		0.16	0.33	0.44	0.04		0.22	0.09
PSI- BLAST	3.60E-51	3.60E-51	7.20E-51	7.20E-51	5.40E-43	5.20E-73	1.10E-36	1.30E-37
End	378	405	433	461	181	298	274	443
Start AA	296	324	352	380	. 66	126	129	297
Chain ID	ပ	υ	v	ပ	ວ	⋖	٧	A
PDB ID	1me y	Ime y	Ime y	Ime y	Ime y	1466	磊	1476
SEQ NO.	277	277	277	277	277	277	772	277

				-				_					_					_					_		_					
POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX ATB ANSCRIPTION	REGULATION/DNA), RNA	POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION	REGULATION/DNA) COMPLEX (TRANSCRIPTION)	REGULATION/DNA), RNA	POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION	REGULATION/DNA) YING-YANG 1;	IKANSCRIPTION INITIATION,	FINGER PROTEIN DNA-PROTEIN	RECOGNITION, 3 COMPLEX	(TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA) YING-YANG 1;	TRANSCRIPTION INITIATION,	INITIATOR ELEMENT, YYI, ZINC 2	PINGER PROTEIN, UNA-PROTEIN	(TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA) YING-YANG 1;	TRANSCRIPTION INITIATION,	INITIATOR ELEMENT, YYI, ZINC 2	FINGER PROTEIN, DNA-PROTEIN	KECOGNITION, 3 COMPLEX (TB ANSCRIPTION BEGIN ATION/DMA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA) YING-YANG 1;	TRANSCRIPTION INITIATION,	INITIATOR ELEMENT, YY1, ZINC 2	FINGER PROTEIN, DNA-PROTEIN RECOGNITION. 3 COMPLEX
	TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, E.	CRAIN. B, C, E, F;		TFIIIA; CHAIN: A, D; 5S	RIBOSOMAL RNA GENE;			YY1; CHAIN: C; ADENO-	ASSOCIATED VIRUS PS	CHAM: A B:	(7 to 1)			YYI; CHAIN: C; ADENO-	ASSOCIATED VIRUS P5	INITIATOR ELEMENT DNA;	CHAIN: A, B;			YY1; CHAIN: C; ADENO.	ASSOCIATED VIRUS P5	INITIATOR ELEMENT DNA;	CHAIN: A, B;			YY I: CHAIN: C: ADENO-	ASSOCIATED VIRUS PS	INITIATOR ELEMENT DNA;	CHAIN: A, B;	
	96.0			0.15				0.87						-						1						_				
	0.15			-0.27				0.12						0.32						0.2						0.3				
	1.80E-36			3.60E-33				3.90E-42						1.80E-32						6.50E-52						1.30E-53				
	461			218				509						237						237						366				
	325			72				120						131						133					_	154				
	4			٧				ပ				-		U						O						U				
	<u>1</u>			1466				lubd					\dashv	_		_				lubd						lubd		_		
	772			772				772						277						772		_				772				
		146 A 325 461 1.80E-36 0.15 0.96 TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE;	1tf6 A 32.5 461 1.80E-36 0.15 0.96 TFIIIA; CHAIN: A, D; SS RIBOSOMAL RNA GENE; CHAIN: B, C, E, F; CHAIN: B, C, E, F;	146 A 325 461 1.80E-36 0.15 0.96 TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	1tf6 A 32.5 461 1.80E-36 0.15 0.96 TFIIIA; CHAIN: A, D; SS RIBOSOMAL RNA GENE; CHAIN: B, C, E, F; CHAIN: B, C, E, F; 1tf6 A 72 218 3.60E-33 -0.27 0.15 TFIIIA; CHAIN: A, D; SS	14f6 A 32.5 461 1.80E-36 0.15 0.96 TFIIA; CHAIN: A, D; SS RIBOSOMAL RNA GENE; CHAIN: B, C, E, F; CHAIN: B, C, E, F; 14f6 A 72 218 3.60E-33 -0.27 0.15 RIBOSOMAL RNA GENE; 14 A N: R D: 218 3.60E-33 -0.27 0.15 RIBOSOMAL RNA GENE;	1tf6 A 32.5 461 1.80E-36 0.15 0.96 TFIIA; CHAIN: A, D; SS RIBOSOMAL RNA GENE; CHAIN: B, C, E, F; CHAIN: B, C, E, F; F; 1tf6 A 72 218 3.60E-33 -0.27 0.15 TFIIIA; CHAIN: A, D; SS 1tf6 A 72 218 3.60E-33 -0.27 0.15 RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	1tf6 A 325 461 1.80E-36 0.15 0.96 TFIIIA; CHAIN: A, D; SS RIBOSOMAL RNA GENE; CHAIN: B, C, E, F; 1tf6 A 72 218 3.60E-33 -0.27 0.15 TFIIIA; CHAIN: A, D; SS RIBOSOMAL RNA GENE; CHAIN: B, C, E, F; CHAIN: B, C, E, F;	1tf6 A 32.5 461 1.80E-36 0.15 0.96 ITFIIIA; CHAIN: A, D; SS RIBOSOMAL RNA GENE; CHAIN: B, C, E, F; CHAIN: B, C, E, F; F; 1tf6 A 72 218 3.60E-33 -0.27 0.15 TFIIIA; CHAIN: A, D; SS 1ubd C 120 209 3.90E-42 0.12 0.87 YYI; CHAIN: C; ADENO-	1tf6 A 325 461 1.80E-36 0.15 0.96 ITFIIIA; CHAIN: A, D; SS RBOSOMAL RNA GENE; CHAIN: B, C, E, F; CHAIN: B, C, E, F; 1tf6 A 72 218 3.60E-33 -0.27 0.15 TFIIIA; CHAIN: A, D; SS 1ubd C 120 209 3.90E-42 0.12 0.87 YY1; CHAIN: C, ADENO-ASSOCIATED VIRUS P5	1tf6 A 325 461 1.80E-36 0.15 0.96 ITFIIIA; CHAIN: A, D; SS RBOSOMAL RNA GENE; CHAIN: B, C, E, F; CHAIN: B, C, E, F; 1tf6 A 72 218 3.60E-33 -0.27 0.15 TFIIIA; CHAIN: A, D; SS 1ubd C 120 209 3.90E-42 0.12 0.87 YYI; CHAIN: C, ADENO-ALAND: A, B. Iubd C 120 3.90E-42 0.12 0.87 YYI; CHAIN: C, ADENO-ALAND: A, B.	1tf6 A 32.5 461 1.80E-36 0.15 0.96 ITFIIIA; CHAIN: A, D; SS 1tf6 A 72 218 3.60E-33 -0.27 0.15 TFIIIA; CHAIN: A, D; SS 1ubd C 120 209 3.90E-42 0.12 0.87 YYYI; CHAIN: C; ADENO-ASSOCIATED VIRUS PS Iubd C 120 3.90E-42 0.12 0.87 YYYI; CHAIN: C; ADENO-ASSOCIATED VIRUS PS Iubd C 120 3.90E-42 0.12 0.87 YYYI; CHAIN: C; ADENO-ASSOCIATED VIRUS PS Iubd C 120 3.90E-42 0.12 0.87 YYYI; CHAIN: C, ADENO-ASSOCIATED VIRUS PS Iubd C 120 3.90E-42 0.12 0.87 YYYI; CHAIN: A, B;	1tf6 A 32.5 461 1.80E-36 0.15 0.96 ITFIIIA; CHAIN: A, D; SS 1tf6 A 72 218 3.60E-33 -0.27 0.15 TFIIIA; CHAIN: A, D; SS 1ubd C 120 209 3.90E-42 0.12 0.87 YYI; CHAIN: C; ADENO-ASSOCIATED VIRUS PS Iubd C 120 209 3.90E-42 0.12 0.87 YYYI; CHAIN: C; ADENO-ASSOCIATED VIRUS PS Iubd C 120 209 3.90E-42 0.12 0.87 YYYI; CHAIN: C; ADENO-ASSOCIATED VIRUS PS Iubd C 120 209 3.90E-42 0.12 0.87 YYYI; CHAIN: A, B;	1tf6 A 325 461 1.80E-36 0.15 0.96 ITIIIA; CHAIN: A, D; SS 1tf6 A 72 218 3.60E-33 -0.27 0.15 TFIIIA; CHAIN: A, D; SS 1ubd C 120 209 3.90E-42 0.12 0.87 YYI; CHAIN: C, ADENO-ASSOCIATED VIRUS PS Iubd C 120 209 3.90E-42 0.12 0.87 YYI; CHAIN: C, ADENO-ASSOCIATED VIRUS PS Iubd C 120 209 3.90E-42 0.12 0.87 YYI; CHAIN: C, ADENO-ASSOCIATED VIRUS PS Iubd C 120 209 3.90E-42 0.12 0.87 XYI; CHAIN: A, B;	1tf6 A 32.5 461 1.80E-36 0.15 0.96 ITIIIA; CHAIN: A, D; SS 1tf6 A 72 218 3.60E-33 -0.27 0.15 Itilia; CHAIN: B, C, E, F; 1ubd C 120 209 3.90E-42 0.12 0.87 YYI; CHAIN: C, ADENO-ASSOCIATED VIRUS P5 1ubd C 131 237 1.80E-32 1 YYI; CHAIN: C, ADENO-YYI; CHAIN: C, AD	1tf6 A 325 461 1.80E-36 0.15 0.96 ITTIIA; CHAIN: A, D; SS 1tf6 A 72 218 3.60E-33 -0.27 0.15 TFIIIA; CHAIN: A, D; SS 1ubd C 120 209 3.90E-42 0.12 0.87 YYI; CHAIN: C, ADENO-ASSOCIATED VIRUS PS 1ubd C 131 237 1.80E-32 0.32 1 1ubd C 131 237 1.80E-32 0.12 0.87	1tf6 A 325 461 1.80E-36 0.15 0.96 TFIIIA; CHAIN: A, D; SS 1tf6 A 72 218 3.60E-33 -0.27 0.15 TFIIIA; CHAIN: A, D; SS 1ubd C 120 209 3.90E-42 0.12 0.87 YY1; CHAIN: C; ADENO-ASSOCIATED VIRUS PS 1ubd C 131 237 1.80E-32 0.32 1 YY1; CHAIN: C; ADENO-ASSOCIATED VIRUS PS 1ubd C 131 237 1.80E-32 1 YY1; CHAIN: C; ADENO-ASSOCIATED VIRUS PS 1ubd C 131 237 1.80E-32 1 YY1; CHAIN: C; ADENO-ASSOCIATED VIRUS PS 1ubd C 131 237 1.80E-32 1 XY1; CHAIN: C; ADENO-ASSOCIATED VIRUS PS 1ubd C 131 237 1 XY1; CHAIN: C; ADENO-ASSOCIATED VIRUS PS	1476 A 325 461 1.80E-36 0.15 0.96 TFIIIA; CHAIN: A, D; SS RIBOSOMAL RNA GENE; CHAIN: B, C, E, F; CHAIN: B, C, E, F; CHAIN: B, C, E, F; CHAIN: B, C, E, F; CHAIN: C, ADENO-C, CHAIN: B, C, E, F; CHAIN: C, ADENO-C, CHAIN: C, ADENO-C, CHAIN: C, ADENO-C, CHAIN: C, ADENO-C, CHAIN: C, ADENO-C, CHAIN: C, ADENO-C, CHAIN: C, ADENO-C, CHAIN: C, ADENO-C, CHAIN: C, ADENO-C, CHAIN: C, ADENO-C, CHAIN: C, ADENO-C, CHAIN: C, ADENO-C, CHAIN: C, ADENO-C, CHAIN: C, ADENO-C, CHAIN: C, ADENO-C, CHAIN: C, ADENO-C, CHAIN: A, B; CHAIN: A, B; CHAIN: A, B; CHAIN: A, B;	1466 A 325 461 1.80E-36 0.15 0.96 TFIIIA; CHAIN: A, D; SS RIBOSOMAL RNA GENE; CHAIN: B, C, E, F; CHAIN: B, C, E, F; CHAIN: B, C, E, F; CHAIN: B, C, E, F; CHAIN: B, C, E, F; CHAIN: B, C, E, F; CHAIN: C, ADENO-ASSOCIATED VIRUS P5 Dibb C 120 209 3.90E-42 0.12 0.87 YY1; CHAIN: C, ADENO-ASSOCIATED VIRUS P5 Dibb C 131 237 1.80E-32 0.32 1 XYY1; CHAIN: C, ADENO-ASSOCIATED VIRUS P5 DIVITIATOR ELEMENT DNA; CHAIN: A, B, CHAIN: A, B, CHAIN: A, B, CHAIN: A, B,	1466 A 325 461 1.80E-36 0.15 0.96 RIBOSOMALRIN: A. D.; SS RIBOSOMALRIN: A. D.; SS CHAIN: B, C, E, F; CHAIN: B, C, E, F; CHAIN: B, C, E, F; CHAIN: B, C, E, F; CHAIN: B, C, E, F; CHAIN: B, C, E, F; CHAIN: B, C, E, F; CHAIN: C, ADENO-ASSOCIATED VIRUS PS CHAIN: C, ADENO-ASSOCIATED VIRUS PS CHAIN: C, ADENO-ASSOCIATED VIRUS PS CHAIN: C, ADENO-ASSOCIATED VIRUS PS CHAIN: C, ADENO-ASSOCIATED VIRUS PS CHAIN: C, ADENO-ASSOCIATED VIRUS PS CHAIN: A, B; CHAIN: A,	1466 A 325 461 1.80E-36 0.15 0.96 TFIIIA; CHAIN: A, D; SS RBOSOMAL RNA GENE; CHAIN: B, C, E, F; CHAIN: B, C, E, F; CHAIN: B, C, E, F; CHAIN: B, C, E, F; CHAIN: B, C, E, F; CHAIN: B, C, E, F; CHAIN: B, C, E, F; CHAIN: B, C, E, F; CHAIN: B, C, E, F; CHAIN: B, C, E, F; CHAIN: C, ADENO-ASSOCIATED VIRUS P5 TIUDA C 131 237 1.80E-32 0.32 1 TY1; CHAIN: C, ADENO-ASSOCIATED VIRUS P5 TRITIATOR ELEMENT DNA; CHAIN: A, B; CHAIN: C, ADENO-ASSOCIATED VIRUS P5 TRITIATOR ELEMENT DNA; CHAIN: C, ADENO-ASSOCIATED VIRUS P5 TRITIATOR CHAIN: C, ADENO-ASSOCIATED VIRUS P5 TRITIATOR CHAIN: C, ADENO-ASSOCIATED VIRUS P5 TRITIATOR CHAIN: C, ADENO-ASSOCIATED VIRUS P5 TRITIATOR CHAIN: C, ADENO-ASSOCIATED VIRUS P5 TRITIATOR CHAIN: C, ADENO-ASSOCIATED VIRUS P5 TRITIATOR CHAIN: C, ADENO-ASSOCIATED VIRU	HE A 325 461 1.80E-36 0.15 0.96 TFIIIA; CHAIN: A, D, SS RIBOSOMAL RNA GENE; CHAIN: B, C, E, F; CHAIN: B, C, E, F; CHAIN: B, C, E, F; CHAIN: B, C, E, F; CHAIN: B, C, E, F; CHAIN: B, C, E, F; CHAIN: C, CHAIN: B, C, E, F; CHAIN: C, CHAIN: B, C, E, F; CHAIN: C, CHAIN: B, C, E, F; CHAIN: C, CHAIN: C, ADENO-CHAIN: C, ADE	HE A 325 461 1.80E-36 0.15 0.96 TFIIIA; CHAIN: A, D; SS RIBOSOMAL RNA GENE; CHAIN: B, C, E, F; CHAIN: B, C, E, F; CHAIN: B, C, E, F; CHAIN: B, C, E, F; CHAIN: B, C, E, F; CHAIN: B, C, E, F; CHAIN: CHAIN: CHAIN: A, D; SS CHAIN: C, ADENO-CHAIN: C, ADEN	HE	1466 A 325 461 1.80E-36 0.15 0.96 TFILIA; CHAIN: A, D; SS RIBOSOMAL RNA GENE; CHAIN: B, C, E, F; C 120 201 3.90E-42 0.15 TFILIA; CHAIN: A, D; SS RIBOSOMAL RNA GENE; CHAIN: B, C, E, F; C 120 209 3.90E-42 0.12 0.87 YY1; CHAIN: C; ADENO-ASSOCIATED VIRUS PS NITIATOR ELEMENT DNA; CHAIN: C, ADENO-ASSOCIATED VIRUS PS NITIATOR ELEMENT DNA; CHAIN: C, ADENO-ASSOCIATED VIRUS PS NITIATOR ELEMENT DNA; CHAIN: C, ADENO-ASSOCIATED VIRUS PS NITIATOR ELEMENT DNA; CHAIN: C, ADENO-ASSOCIATED VIRUS PS NITIATOR ELEMENT DNA; CHAIN: C, ADENO-ASSOCIATED VIRUS PS NITIATOR ELEMENT DNA; CHAIN: A, B; RIBARD	146 A 325 461 1.80E-36 0.15 0.96 TFILIA; CHAIN: A, D; SS RIBOSOMAL RNA GENE; CHAIN: B, C, E, F; CHAIN: B, C, E, F; CHAIN: C, E, E, F; CHAIN: C, E, E, F; CHAIN: C, E, E, F; CHAIN: C, E, E, E, E, E, E, E, E, E, E, E, E, E,	1466 A 325 461 1.80E-36 0.15 0.96 TFIIIA; CHAIN: A, D; SS RIBOSOMAL, RNA GENE; CHAIN: B, C, E, F; CHAIN: B, C, E, F; CHAIN: C, C, E, F; CHAIN: C, C, E, F; CHAIN: C, C, C, E, F; CHAIN: C, C, C, E, F; CHAIN: C, C, C, E, F; CHAIN: C, C, C, E, F; CHAIN: C, C, C, E, F; CHAIN: C, C, C, E, F; CHAIN: C, C, C, E, F; CHAIN: C, C, C, E, F; CHAIN: C, C, C, E, F; CHAIN: C, C, C, E, F; C, C, C, C, C, C, C, C, C, C, C, C, C,	1466 A 325 461 180E-36 0.15 0.96 TFILIA; CHAIN: A, D; SS RIBOSOMAL RNA GENE; CHAIN: B, C, E, F; CHAIN: B, C, E, F; CHAIN: C, ADENO 1.90E-42 0.15 0	1476 A 325 461 180E-36 0.15 0.96 TFILIA; CHAIN: A, D; SS RIBOSOMAL RNA GENE; CHAIN: B, C, E, F; CHAIN: C, ADENO-B, C, CHAIN: B, C, E, C, C, C, C, C, C, C, C, C, C, C, C, C,	1476 A 325 461 180E-36 0.15 0.96 TFILIA; CHAIN: A, D; SS RIBOSOMAL RNA GENE; CHAIN: B, C, E, F; CHAIN: B, C, E, F; CHAIN: C, D; SS CHAIN: B, C, E, F; CHAIN: C, D; SS CHAIN: B, C, E, F; CHAIN: C, D; SS CHAIN: C, C, CHAIN: C

PDB annotation	(TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATIONDNA) YING-YANG 1; TRANSCRIPTION INITIATION,	INITIATOR ELEMENT, YY1, ZINC 2	FINGER PROTEIN, DNA-PROTEIN	TRECOGNITION, 3 COMPLEX (TRANSCRIPTION REGIII.ATION/DNA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA) YING-YANG 1;	TRANSCRIPTION INITIATION,	ENICEP PROTEIN DAY PROTEIN	RECOGNITION, 3 COMPLEX	(TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA) YING-YANG 1;	TRANSCRIPTION INITIATION,	INITIATOR ELEMENT, YYI, ZINC 2	PECOCAITTON 2 COMPLEX	RECOGNITION, 3 COMPLEX	COMPLEX (TRANSCRIPTION	REGIII ATION/DNA) YING-YANG I	TRANSCRIPTION INITIATION.	INITIATOR ELEMENT, YY1, ZINC 2	FINGER PROTEIN, DNA-PROTEIN	RECOGNITION, 3 COMPLEX	(TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION PEGIT ATTOMONA) VING YANG 1:	TRANSCRIPTION INITIATION.	INITIATOR ELEMENT, YYI, ZINC 2	FINGER PROTEIN, DNA-PROTEIN	RECOGNITION, 3 COMPLEX	(TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA) YING-YANG 1;	DATE ATOR ELEMENT AND ZELOS	FINGER PROTEIN, DNA-PROTEIN
Coumpound		YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA;	CHAIN: A, B;			YY1; CHAIN: C; ADENO-	ASSOCIATED VIRUS P5	INITIATOR ELEMENT DNA;	CHAIN: A, B;			YY1; CHAIN: C; ADENO-	ASSOCIATED VIRUS PS	INITIATOR ELEMENT DNA;	CHAIN: A, B;			VY1. CHAIN: C. ADENO.	ASSOCIATED VIRUS P5	INITIATOR ELEMENT DNA:	CHAIN: A, B;				YY1; CHAIN: C; ADENO-	INITIATOR ELEMENT DNA:	CHAIN: A, B;				YYI; CHAIN: C; ADENO-	ASSOCIATED VIRUS PS	CHAN: A B.	CHOIN, A, D,
SeqFold score		89.18	•																															
PMF score						0.92	-					96.0						860	2						, -						0.95			
Verify score						-0.04						0.26						0.22							0.16						4.0			
PSI- BLAST		7.80E-55				7.80E-55						6.50E-56						1 60E-35							1.30E-56						5.20E-51			
End		322				349						405						405	:						434						194			
Start AA		212				238				•		294						304							322						350			
Chain		U				U						ပ						C	1						່		-	_			ບ			
PDB CO		1ubd				Jubd						1 pq				_		1ubd							pqn	_				-	Iubd			
SEQ No. 19		772				772						772				_		772						į	7//						77.7			

PDB annotation	RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATIONDNA) YING-YANG I; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YYI, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATIONDNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC PINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEINIDNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/INA)	COMPLEX (DNA-BINDING PROTEINDNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEINDNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEINDNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA-	ENCHALT INCITED TO THE PROPERTY OF THE PROPERT
Coumpound		YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;	ZINC FINGER PROTEIN GL11; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;
SeqFold score						98.73				
PMF		0.92	0.88	_	0.76		0.93	0.84	96.0	1
Verify score		0.32	0.21	0.35	0.16		0.4	0.24	0.35	0.47
PSI- BLAST		1.805-34	2.60E-54	3.60E-33	3.90E-66	1.00E-71	1.00E-71	7.20E-33	6.50E-67	5.40E-34
End		461	239	264	295	351	407	432	461	460
Start AA		360	121	128	128	212	268	304	324	332
Chain ID		o	V	A.	¥	⋖ . :	٧	٧	٧	V
PDB ID		lubd	2gli	2gli	2gli	2gli	2gli	2gli	2gli	2gli
SEQ B NO:		277	772	772	772	772	277	277	772	772

			_				_				7					
PDB annotation	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)		HYDROLASE TETRATRICOPEPTIDE, TRP; HYDROLASE, PHOSPHATASE, PROTEIN-PROTEIN INTERACTIONS, TPR, 2 SUPER-HELIX, X-RAY STRUCTURE	CHAPERONE HOP, TPR-DOMAIN, PEPTIDE-COMPLEX, HELICAL REPEAT, HSP90, 2 PROTEIN BINDING	CHAPERONE HOP, TPR-DOMAIN, PEPTIDE-COMPLEX, HELICAL REPEAT, HSC70, 2 HSP70, PROTEIN BINDING	SIGNALING PROTEIN PEROXISMORE	RECEPTOR 1, PTS1-BP, PEROXIN-5,	TETRATRICOPEPTIDE REPEAT, TPR. 2	HELICAL REPEAT		AMIDOTRANSFERASE AMIDOTRANSFERASE, THIOESTER	AMIDOTRANSFERASE AMIDOTRANSFERASE, THIOESTER	LYASE AIRC, PURK; ATP-GRASP, CARBOXYPHOSPHATE, PURINE BIOSYNTHESIS, LYASE	LIGASE ATP-GRASP, CARBOXYLASE, BIOTIN-DEPENDENT	LIGASE LMDDL2; ATP-BINDING. GRASP MOTIF FOR ATP.
Coumpound	ZINC FINGER PROTEIN GLI1; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;		SERNE/THREONINE PROTEIN PHOSPHATASE 5; CHAIN: NULL;	TPR2A-DOMAIN OF HOP; CHAIN: A; HSP90-PEPTIDE MEEVD; CHAIN: B;	TPR1-DOMAIN OF HOP; CHAIN: A, B; HSC70- PEPTIDE; CHAIN: C, D;	PEROXISOMAL TARGETING	SIGNAL I RECEPTOR;	CONTAINING PEPTIDE	CHAIN: C, D;		CARBAMOYL PHOSPHATE SYNTHETASE; CHAIN: A, B, C, D, E, F, G, H;	CARBAMOYL PHOSPHATE SYNTHETASE; CHAIN: A, B, C, D, E, F, G, H;	NS- CARBOXYAMINOIMIDAZOL E RIBONUCLEOTIDE CHAIN: A:	BIOTIN CARBOXYLASE; CHAIN: A. B:	D-ALANINE:D-LACTATE LIGASE; CHAIN: A, B;
SeqFold score																
PMF	-0.05	0.15		0.77	99.0	0.74	66 0					6.0	96:0	-0.11	0.54	0.45
Verify score	0.07	-0.01		0.09	0.14	0.12	0.47	:				0.14	0.49	0.07	0.32	0.07
PSI- BLAST	9.00E-28	5.40E-31		1.30E-08	9.10E-08	6.50E-10	6.50E-09					1.80E-64	1.10E-39	1.80E-16	1.30E-86	9.00E-37
End	180	211		180	180	180	179					860	892	828	965	858
Start AA	20	71		105	105	105	105	}				454	529	532	534	531
Chain ID	∢	٧			¥	۲	▼					٧	A	¥	A	A
PDB ID	2gli	2gli		la17	lefr	1elw	lfch					1в9х	1а9х		ldvl	lchi
SE SE	277	772		577	773	773	773					774	774	774	774	774

PDB annotation	LIGASE LMDDL2; ATP-BINDING. GRASP MOTIF FOR ATP.	LIGASE SCS-ALPHA; SCS-BETA; LIGASE, GTP-SPECIFIC 1;	A TRANSFERASE TRANSFORMYLASE, PURINE BIOSYNTHESIS, ATP-GRASP	A TRANSFERASE TRANSFORMYLASE, PURINE BIOSYNTHESIS, ATP-GRASP	LIGASE PURD GEN PRODUCT; GAR- SYN, GLYCINAMIDE RIBONUCLEOTIDE SYNTHETASE, ATP-GRASP, 2 PURINE DE NOVO BIOSYNTHETIC PATHWAY, SUBSTRATE CHANNELING	LIGASE DD-LIGASE, DDLB; GLYCOGEN PHOSPHORYLASE, LIGASE, CELL WALL, PEPTIDOGLYCAN 2 SYNTHESIS, VANCOMYCIN, ADP BINDING	LIGASE SCS; SCRITRIC ACID CYCLE, HETEROTETRAMER, LIGASE	TRANSLATION PROTEIN-PROTEIN COMPLEX	INTEGRIN INTEGRIN, CELL ADHESION, GLYCOPROTEIN	INTEGRIN INTEGRIN, CELL ADHESION, GLYCOPROTEIN	INTEGERAL INTEGERAL COLI
Соитроипа	D-ALANINE:D-LACTATE LIGASE; CHAIN: A, B;	SUCCINYL-COA SYNTHETASE, ALPHA CHAIN: CHAIN, A; SUCCINYL-COA SYNTHETASE, BETA CHAIN; CHAIN: B;	PHOSPHORIBOSYLGLYCINA MIDE FORMYLTRANSFERASE 2; CHAIN: A. B:	PHOSPHORIBOSYLGLYCINA MIDE FORMYLTRANSFERASE 2; CHAIN: A. B:	GLYCINAMIDE RIBONUCLEOTIDE SYNTHETASE; CHAIN: A;	D-ALAUD. ALA LIGASE; CHAIN: NULL;	SUCCINYL-COA LIGASE; CHAIN: A, D; SUCCINYL- COA LIGASE; CHAIN: B, E;	ELONGATION FACTOR EEFIA; CHAIN: A; ELONGATION FACTOR EEFIBA; CHAIN: B;	INTEGRIN ALPHA 2 BETA; CHAIN: A. B:	INTEGRIN ALPHA 2 BETA; CHAIN: A, B;	NITECONI AT DHA 2 DETA.
SeqFold										84.82	
PMF	0.63	0.07	0.99	0.11	0.37	0.18	0.07	-	_		-
Verify score	0.22	-0.04	0.51	0.21	0.5	-0.07	0.04	1.05	99.0		0 74
PSI- BLAST	1.80E-36	7.20E-22	6.50E-30	1.40E-39	3.60E-54	1.80E-43	1.80E-27	0	9.00E-28	9.00E-28	7.20E-28
End	858	854	953	098	868	. 859	844	443	313	316	504
Start AA	531	959	533	534	531	530	929	2	119	120	321
Chain ID	æ	Ø	4	¥	¥		В	4	V.	<	4
PDB ID	1ehi	1euc	leyz	leyz	1gso	liow	2scu	1160	laox	laox	laox
SEQ NO.	774	774	774	774	774	774	774	775	776	176	176

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PDB annotation	CYTOSKELETON	CELL ADHESION PROTEIN A-DOMAIN INTEGRIN, CELL ADHESION PROTEIN, GLYCOPROTEIN, EXTRACELLULAR 2 MATRIX, CYTOSKELETON	CELL ADHESION PROTEIN A-DOMAIN	INTEGRIN, CELL ADHESION PROTEIN CI VCOBROTEIN	EXTRACELLULAR 2 MATRIX, CYTOSKEL FTON	CELL ADHESION PROTEIN A-DOMAIN	INTEGRIN, CELL ADHESION PROTEIN OF YCOPROTEIN	EXTRACELLULAR 2 MATRIX, CYTOSKELETON	CELL ADHESION PROTEIN A-DOMAIN	INTEGRIN, CELL ADRESION	EXTRACELLULAR 2 MATRIX,	CYTOSKELETON	CELL ADHESION LFA-1, ALPHA-	L\BETA-2 INTEGRIN, A-DOMAIN; ILFA 8	CELL ADHESION LFA-1, ALPHA-	L\BETA-2 INTEGRIN; A-DOMAIN; 1LFA 8	CELL ADHESION LFA-1, ALPHA- L\BETA-2 INTEGRIN, A-DOMAIN; LLFA 8	CELL ADHESION INTEGRIN, CELL ADHESION	CELL ADHESION INTEGRIN, CELL ADHESION		SCAFFOLD PROTEIN SCAFFOLD	PROTEIN, PP2A, PHOSPHORYLATION, HEAT REPEAT	TRANSPORT PROTEIN SERINE-RICH
Coumpound		INTEGRIN; CHAIN: NULL;	INTEGRIN CHAIN NIII.I.			INTEGRIN; CHAIN: NULL;			INTEGRIN; CHAIN: NULL;				CD11A; 1LFA 5 CHAIN: A. B;	ILFA 6	CD11A; 1LFA 5 CHAIN: A, B;	1LFA 6	CD11A; 1LFA 5 CHAIN: A, B; 1LFA 6	ALPHAI BETAI INTEGRIN; CHAIN: A; ALPHAI BETAI INTEGRIN: CHAIN: B:	ALPHAI BETAI INTEGRIN; CHAIN: A; ALPHAI BETAI	INTEGKIN; CHAIN: B;	PROTEIN PHOSPHATASE	PP2A; CHAIN: A, B;	KARYOPHERIN ALPHA;
SegFold score			14 71	•											90.24								
PMF		_				_			-				-				1	-	1		0.17		-
Verify score		0.62				0.57			0.54				0.42				0.85	0.7	1.09		0.02		0.52
PSI- BLAST		5.40E-27	0 105 44			9.10E-44			9.00E-26				3.60E-23		3.60E-24		3.60E-24	7.20E-28	1.60E-29		3.60E-33		3.60E-49
End		307	700	3		497			498				268		501		503	309	499		343		322
Start		124	324	1		325			326				123		323		326	122	324		_		_
Chain					- 								<		A		4	V	∢		¥		Y
PDB ID		lido	9:5	3		lido			1ido				11fa		1lfa		1lfa	14c5	Iqc5		163u		lee4
SEQ EQ		776	200	3		776			911				176		776		176	776	776		111		111

PDB annotation	RNA POLYMERASE I SUPPRESSOR PROTEIN; ARM REPEAT	TRANSPORT PROTEIN SERINE-RICH RNA POLYMERASE I SUPPRESSOR PROTEIN; ARM REPEAT	NUCLEAR IMPORT RECEPTOR KARYOPHERIN ALPHA; NUCLEAR IMPORT RECEPTOR, NUCLEAR LOCALIZATION SIGNAL, 2 ARMADILLO REPEATS, AUTONHIBITION, INTRASTERIC REGULATION	NUCLEAR IMPORT RECEPTOR KARYOPHERIN ALPHA; NUCLEAR IMPORT RECEPTOR, NUCLEAR LOCALIZATION SIGNAL, 2 ARMADILLO REPEATS, AUTOINHIBITION, INTRASTERIC REGULATION	STRUCTURAL PROTEIN ARMADILLO REPEAT, BETA-CATENIN, STRUCTURAL PROTEIN	ARMADILLO REPEAT ARMADILLO REPEAT, BETA-CATENIN, CYTOSKELETON	ARMADILLO REPEAT ARMADILLO REPEAT, BETA-CATENIN, CYTOSKELETON	ENDOCYTOSIS/EXOCYTOSIS SYNAPTOTAGMIN, C2-DOMAIN, EXOCYTOSIS, NEUROTRANSMITTER 2 RELEASE, ENDOCYTOSIS/EXOCYTOSIS	ENDOCYTOSIS/EXOCYTOSIS SYNAPTOTAGMIN, C2-DOMAIN, EXOCYTOSIS, NEUROTRANSMITTER 2 RELEASE, ENDOCYTOSIS/EXOCYTOSIS
Coumpound	CHAIN: A, B; MYC PROTO- ONCOGENE PROTEIN; CHAIN: C, D, E, F;	KARYOPHERIN ALPHA; CHAIN: A, B; MYC PROTO- ONCOGENE PROTEIN; CHAIN: C, D, E, F;	IMPORTIN ALPHA; CHAIN: A;	IMPORTIN ALPHA; CHAIN: A;	BETA-CATENIN; CHAIN: NULL;	BETA-CATENIN; CHAIN: NULL;	BETA-CATENIN; CHAIN: NULL;	SYNAPTOTAGMIN I; CHAIN: A;	SYNAPTOTAGMIN I; CHAIN: A;
SeqFold score			52.47						
PMF score		0.17		-	0.98	0.94	0.75		1
Verify score		0.03		0.55	0.26	0.3	0.21	0.41	0.59
PSI- BLAST		5.40E-10	1.40E-51	1.40E-51	3.60E-39	1.30E-29	5.40E-31	2.60E-39	3.60E-24
End AA		347	344	343	347	336	347	265	264
Start AA		233	-	۶	. 26	-	89	140	143
Chain ID		∢	∢	V				4	∢
PDB CI		1ce4	lial	lial	2bct	3bct	3bct	1byn	1byn
SEQ D NO:		111	<i>TTT</i>	777	777	777	777	779	61.1

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PDB annotation	ENDOCYTOSIS/EXOCYTOSIS BETA SANDWICH, CALCIUM ION, C2 DOMAIN	ENDOCYTOSIS/EXOCYTOSIS BETA SANDWICH, CALCIUM ION, C2 DOMAIN					COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	CONTRACTILE LIM DOMAIN, CRP, NMR, MUSCLE DIFFERENTIATION. CONTRACTILE	SIGNALLING PROTEIN BINDING PROTEIN, CYTOKINE, SIGNALLING PROTEIN	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX
Coumpound	SYNAPTOTAGMIN III; CHAIN: A;	SYNAPTOTAGMIN III; CHAIN: A;	CALCIUMPHOSPHOLIPID BINDING PROTEIN SYNAPTOTAGMIN I (FIRST C2 DOMAIN) (CALB) IRSY 3	CALCIUM/PHOSPHOLIPID BINDING PROTEIN SYNAPTOTAGMIN I (FIRST C2 DOMAIN) (CALB) IRSY 3	CALCIUMPHOSPHOLIPID BINDING PROTEIN SYNAPTOTAGMIN I (FIRST C2 DOMAIN) (CALB) IRSY 3		QGSR ZINC FINGER PEPTIDE, CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE, CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	CRP1; CHAIN: A;	TUMOR NECROSIS FACTOR RECEPTOR: CHAIN: A, B:	DNA; CHAIN: A, B, D, E; CONSENSUS ZNC FINGER PROTEIN; CHAIN: C, F, G;
SeqFold score		_	130.23				70.71			58.15	57.16	
PMF	_	1		_	-			0.03	90.0	,		-
Verify score	0.51	0.43		0.35	0.56			-0.31	-0.2			0.32
PSI- BLAST	2.60E-73	9.00E-55	2.60E-41	2.60E-41	3.60E-24		7.20E-31	7.20E-22	3.60E-26	7.80E-13	0.00026	3.60E-50
End	376	387	266	264	264		227	16	169	277	186	197
Start AA	140	143	136	140	143		145	19	95	88	01	116
Chain	V	A					<	4	V	∢	¥	ပ
PDB	1dqv	ldqv	Irsy	1139	Irsy		lalh	lalh	lath	168t	lext	Ime y
SEQ PO PO PO PO PO PO PO PO PO PO PO PO PO	977	779		977	977		783	783	783	783	783	783

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PDB annotation	(ZINC FINGER/DNA)	COMPLEX (ZINC FINGENDNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2	CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC	FINGER, PROTEIN-DNA	CRYSTAL STRUCTURE, COMPLEX	(ZINC FINGER/DNA)	COMPLEX (ZINC FINGENDNA) ZINC	FINGER, PROTEIN-DNA INTERACTION PROTEIN DESIGN 2	CRYSTAL STRUCTURE, COMPLEX	(ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC	FINGER, PROTEIN-DNA	INTERACTION, PROTEIN DESIGN, 2	CRYSTAL STRUCTURE, COMPLEX	COMPLEX (ZINC FINGER/DNA) ZINC	FINGER, PROTEIN-DNA	INTERACTION, PROTEIN DESIGN, 2	CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC	FINGER, PROTEIN-DNA	INTERACTION. PROTEIN DESIGN, 2	CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC	FINGER, PROTEIN-DNA	INTERACTION, PROTEIN DESIGN, 2	CRYSTAL STRUCTURE, COMPLEX	(ZINC FINGER/DNA)	COMPLEX (TRANSCRIPTION	KEGULATION/DNA) COMPLEX	REGULATION/DNA), RNA	POLYMERASE III, 2 TRANSCRIPTION
Coumpound		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;		DNA; CHAIN: A, B, D, E,	CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;		DNA; CHAIN: A, B, D, E;	CONSENSOS ZINC FINGER PROTEIN: CHAIN: D. R. G.	INCIDENT, CIETTIN, C, I, C,		DNA; CHAIN: A, B, D, E;	CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;		DNA: CHAIN: A. B. D. E.	CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;		DNA; CHAIN: A, B, D, E;	CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;		DNA; CHAIN: A, B, D, E;	CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;			TFIIIA; CHAIN: A, D; 5S	CHAIN: B.C.E.F.		
SeqFold score								92.4																					87.93			
PMF		-		_								0.03				_				0.03				0.74								
Verify score		0.54		0.47								-0.15				0.43			ü	-0.51				90.0								
PSI- BLAST		3.60E-50		1.30E-50				1.30E-50				3.60E-39				1.80E-46				5.40E-42				1.10E-45					1.30E-58			
End AA		225		253				254				16				276				141				169					275			
Start AA		144		172				172				18				200				89				94					911			
Chain ED in		ပ		ပ				ပ				၁				ပ				O				ပ					٧			
PDB ID		, Ime		Ime	>			Ime	>			Ime	>			Ime	^			1mc	>			- Inc	>			1	9#1			
SEQ No B		783		783				783				783				783				783				783					783			

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PDB annotation	INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION,	INITIATOR ELEMENT, YY1, ZINC 2	FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX	(TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA) YING-YANG 1;	I KANSCELL TION INTITATION,	FINGER PROTEIN, DNA-PROTEIN	RECOGNITION, 3 COMPLEX	(TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA) YING-YANG I;	INTITATOR ELEMENT, YY1, ZINC 2	FINGER PROTEIN, DNA-PROTEIN	RECOGNITION, 3 COMPLEX	(TRANSCRIPTION REGULATION/UNA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA) YING-YANG I;	IRANSCRIPTION INITIATION,	FINGER PROTEIN DNA-PROTEIN	RECOGNITION, 3 COMPLEX	(TRANSCRIPTION REGULATION/DNA)	COMPLEX (DNA-BINDING	PROTEIN/DINA) FIVE-FINGER GEI, GEI,	ZINC FINGER, COMPLEX (DNA- RINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING	PROTEIN/DNA) FIVE-FINGER GLI; GLI,	ZINC FINGER, COMPLEX (DNA-	BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING	PROTEIN/DINA) FIVE-FINGER GLI; GLI,	BINDING PROTEIN/DNA)
Coumpound		YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS PS INITIATOR FI EMENT DIVA:	CHAIN: A, B;			YY1; CHAIN: C; ADENO-	ASSOCIATED VIRUS P5	CHAN: A R.	(T (1) (1) (1)	•		YY1; CHAIN: C; ADENO-	ASSOCIATED VIRUS PS	CHAIN A B.	,			YY1; CHAIN: C; ADENO-	ASSOCIATED VIRUS P5	INITIATOR ELEMENT DNA;	CHAIN: A, B;			ZINC FINGER PROTEIN GLII;	CHAIN: A; DNA; CHAIN: C,	<u>:</u>	ZINC FINGER PROTEIN GLII:	CHAIN: A; DNA; CHAIN: C,	Ö		ZINC FINGER PROTEIN GLII;	CHAIN: A; DNA; CHAIN: C,	á
SeqFold score		85.55																						86.72									
PMF						0.87						1						0.46									-	•			6.0		
Verify			`			0.1						-0.02						-0.12								-	0.14				0.23		
PSI- BLAST		3.90E-51				9.10E-47						3.90E-51						3.60E-32						3.90E-59			3 OOF - 59	70.70			1.30E-56		•
End		226				225						254						197						255			255	}			267		•
Start		116				121						149						70						116			117	:			145		
Chain ID		ပ				၁						ပ						O						٧			4	:			Ą		
PDB ID		lubd	•			lubd						lubd						1ubd						2gli			Joli	19			2gli		
SEQ.	5	783				783						783				_		783						783			783	3			783		

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PDB annotation	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)		TRANSFERASE ATK, AMGXI, BPK; TYROSINE KINASE, X-LINKED AGAMMAGLOBULINEMIA, XLA, BTK, SH3 2 DOMAIN, TRANSFERASE	COMPLEX (ADAPTOR PROTEIN/PEPTIDE) ASH, GROWTH FACTOR RECEPTOR-BOUND PROTEIN 2; COMPLEX (ADAPTOR PROTEIN/PEPTIDE), SH3 DOMAIN, 2 GUANINE-NUCLEOTIDE RELEASING FACTOR	COMPLEX (TRANSFERASE/PEPTIDE) COMPLEX (TRANSFERASE/PEPTIDE), SIGNAL TRANSDUCTION, 2 SH3 DOMAIN	COMPLEX (SIGNAL TRANSDUCTION/PEPTIDE) COMPLEX (SIGNAL TRANSDUCTION/PEPTIDE), SH3 DOMAIN	SIGNAL, TRANSDUCTION ADAPTOR SH2, SH3 IGRI 14	CIRCULAR PERMUTANT PWT; CIRCULAR PERMUTANT, SH3 DOMAIN, CYTOSKELETON	CYTOSKELETON CYTOSKELETON, MEMBRANE, SH3 DOMAIN	SIGNAL TRANSDUCTION PROTEIN SRC-HOMOLOGY 3 (SH3) DOMAIN, PEPTIDE-BINDING PROTEIN, ISEM 18 2 GUANINE NUCLEOTIDE EXCHANGE FACTOR ISEM 19		ENDOCYTOSIS/EXOCYTOSIS NSECI; PROTEIN-PROTEIN COMPLEX, MULTI-SUBUNIT
Coumpound	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;		BRUTON'S TYROSINE KINASE; CHAIN: NULL;	GRB2; CHAIN: A; SOS; CHAIN: B;	ABL TYROSINE KINASE; CHAIN: A, C, E, G; PEPTIDE P41; CHAIN: B, D, F, H;	GRB2; CHAIN: A; SOS-1; CHAIN: B;	GROWTH FACTOR BOUND PROTEIN 2; 1GRI 5 CHAIN: A, B; 1GRI 6	ALPHA SPECTRIN; CHAIN: NULL;	ALPHA II SPECTRIN; CHAIN: A;	SEM-5: ISEM 3 CHAIN: A, B; ISEM 5 10-RESIDUE PROLINE-RICH PEPTIDE FROM MSOS ISEM 8 CHAIN: C, D ISEM 10		SYNTAXIN BINDING PROTEIN 1; CHAIN: A; SYNTAXIN 1A; CHAIN: B;
SeqFold score												
PMF	0.1		9.0	0.83	0.25	0.87	0.12	0.59	0.71	0.59		0.01
Verify score	-0.43		0.42	0.35	0.08	0.4	0.12	0.2	-0.25	0.21		-0.67
PSI- BLAST	3.60E-26		1.00E-09	2.60E-11	3.90E-10	9.10E-12	1.80E-09	2.60E-11	1.30E-10	5.40E-11		1.00E-05
End	140		403	402	403	403	400	403	403	400	_	281
Start	26		340	348	350	348	343	344	349	349		160
Chain ID	¥			∢	4	¥	4		4	∢		æ
EDB TD	2gli		law w	laze	1bbz	1gbq	lgri	lpwt	lqk w	lsem		ldn1
SEQ PO PO PO PO PO PO PO PO PO PO PO PO PO	783		784	784	784	784	784	784	784	784		785

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PDB annotation	HAT OBED ON TO A SE	BROMOPEROXIDASE L, HALOPEROXIDASE L; HALOPEROXIDASE, OXIDOREDUCTASE	HALOPEROXIDASE CHLOROPEROXIDASE A1, HALOPEROXIDASE A1; HALOPEROXIDASE, OXIDOREDUCTASE	HALOPEROXIDASE HALOPEROXIDASE F; HALOPEROXIDASE, OXIDOREDUCTASE, PROPIONATE COMPLEX	HALOPEROXIDASE HALOPEROXIDASE A2, CHLOROPEROXIDASE A2, HALOPEROXIDASE, OXIDOREDUCTASE, PEROXIDASE, ALPHABETA 2 HYDROLASE FOLD, MUTANT M991	HYDROLASE BPHD; HYDROLASE, PCB DEGRADATION	HYDROLASE A/B HYDROLASE FOLD, DEHALOGENASE I-S BOND	HYDROLASE TRIACYLGLYCEROL- HYDROLASE, X-RAY CRYSTALLOGRAFHY. PSEUDOMONADACEAE, OXYANION, CIS-PEPTIDE, HYDROLASE	HYDROLASE HYDROLASE, ALPHA/BETA HYDROLASE FOLD, EPOXIDE DEGRADATION, 2 EPICHLOROHYDRIN	HYDROLASE HOMODIMER, ALPHA/BETA HYDROLASE FOLD, DISUBSTITUTED UREA 2 INHIBITOR
Coumpound	T OF LATTER A POP 1	CHLOROPEROXIDANE L; CHAIN: A, B, C;	BROMOPEROXIDASE A1; CHAIN: NULL;	CHLOROPEROXIDASE F; CHAIN: NULL;	BROMOPEROXIDASE A2; CHAIN: NULL;	2-HYDROXY-6-OXO-6- PHENYLHEXA-2,4- DIENOATE CHAIN: A;	HALOALKANE DEHALOGENASE; 1- CHLOROHEXANE CHAIN: A;	TRIACYLGLYCEROL HYDROLASE; CHAIN: NULL;	SOLUBLE EPOXIDE HYDROLASE; CHAIN: A. B, C. D;	EPOXIDE HYDROLASE; CHAIN: A, B;
SeqFold score							-			
PMF score		0.07	0.05	0.22	0.28	0.87	0.99	0.03	0.52	0.15
Verify score		-0.21	-0.13	-0.12	0.14	-0.02	0.35	0.09	0.23	0
PSI- BLAST		5.40E-45	1.10E-38	7.20E-39	1.30E-39	5.40E-41	1.80E-40	3.60E-11	5.40E-38	1.10E-37
End		-258	257	258	. 258	247	257	203	256	258
Start			6	7	7	17	13	25	=	41
Chain		∀				<	4		V.	4
PDB UD		1a88	1a8q	1a8s	1brt	1c4x	1cqw	levl	lehy	leki
SEQ EQ		786	786	786	786	786	786	786	786	786

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PDB annotation	HYDROLASE HOMODIMER, ALPHA/BETA HYDROLASE FOLD, DISUBSTITUTED UREA 2 INHIBITOR	HYDROLASE LIPASE		HYDROLASE PSEUDOMONADACEAE, CIS-PEPTIDE, CLOSED CONFORMATION, 2 HYDROLASE, LID	HYDROLASE ALPHA BEIA HYDROLASE FOLD, PROLINE, PROLYL AMINOPEPTIDASE, 2 SERRATIA, IMINOPEPTIDASE	LIPASE LIPASE; LIPASE, HYDKULASE, PSEUDOMONADACEAE, COVALENT INTERMEDIATE, 2 TRIGLYCERIDE ANALOGUE, ENANTIOSELECTIVITY	A TACOLO LA CALLED SA COLO	SIGNAL TRANSDUCTION SIGNAL TRANSDUCTION, SOS, PLECKSTRIN HOMOLOGY (PH) DOMAIN	SIGNAL TRANSDUCTION PROTEIN	TRANSPORT PROTEIN RHO-GTPASE EXCHANGE FACTOR, TRANSPORT PROTEIN	TRANSPORT PROTEIN RHO-GTPASE EXCHANGE FACTOR, TRANSPORT PROTEIN	GENE REGULATION SON OF SEVENLESS PROTEIN; GUANINE NUCLEOTIDE EXCHANGE FACTOR, GENE REGULATION
Coumpound	EPOXIDE HYDROLASE; CHAIN: A, B;	LIPASE, GASTRIC; CHAIN: A, B;	HYDROLASE(CARBOXYLIC ESTERASE) LIPASE (E.C.3.1.13) COMPLEXED WITH COLIPASE AND INHIBITED ILPB 3 BY UNDECANE PHOSPHONATE METHYL ESTER (TWO CONFORMATIONS) ILPB 4	TRIACYLGLYCEROL HYDROLASE; CHAIN: D; TRIACYLGLYCEROL HYDROLASE; CHAIN: E;	PROLYL AMINOPEPTIDASE; CHAIN: A;	TRIACYL-GLYCEROL- HYDROLASE; CHAIN: D, E;		SOS1; CHAIN: NULL;	BETA-SPECTRIN; IBTN 4 CHAIN: NULL; IBTN 5	PIX; CHAIN: A;	PIX; CHAIN: A;	HUMAN SOS 1; CHAIN: A;
SeqFold score												
PMF	0.13	0	0	-0.03	0	0.01		0.49	_	0.89	9.0	0.96
Verify	-0.11	-0.32	-0.29	0.1	-0.16	-0.09		0.32	0.39	-0.25	-0.46	-0.1
PSI- BLAST	1.10E-37	1.60E-07	1.40E-06	3.60E-11	1.60E-28	1.60E-11		1.20E-23	9.00E-09	2.60E-41	7.20E-23	1.80E-16
End	258	121	149	203	242	161		576	819	463	462	570
Start	14	٥.	30	25	∞	25		462	728	261	267	261
Chain	В	V	Ф	D	∢	Q				4	4	A
PDB	lek1	Ihlg	11pb	1qge	19tr	4lip		lawe	1btn	1by1	1by1	1dbh
SEQ	NO:	786	786	786	786	786		788	788	788	788	788

PDB annotation	GENE REGULATION SON OF SEVENLESS PROTEIN; GUANINE NUCLEOTIDE EXCHANGE FACTOR, GENE REGULATION	CYTOSKELETON	TRANSFERASE HRS; HRS, VHS, FYVE, ZINC FINGER, SUPERHELJX	SIGNALING PROTEIN 11 ALPHA- HELICES	SIGNALING PROTEIN DAPP'I, PHISH, BAM32; PLECKSTRIN, 3-PLOSPHOINOSITIDES, INOSITOL TETRAKISPHOSPHATE 2 SIGNAL TRANSDUCTION PROTEIN, ADAPTOR PROTEIN	SIGNALING PROTEIN DAPPI, PHISH, BAM32; PLECKSTRIN, 3- PHOSPHONOSTIDES, INOSITOL TETRAKISPHOSPHATE 2 SIGNAL TRANSDUCTION PROTEIN, ADAPTOR PROTEIN	SIGNALING PROTEIN DAPPI, PHISH, BAM32; PLECKSTRIN, 3- PHOSPHOINSTIIDES, INOSITOL TETRAKISPHOSPHATE 2 SIGNAL TRANSDUCTION PROTEIN, ADAPTOR PROTEIN	SIGNALING PROTEIN DAPP1, PHISH, BAM32; PLECKSTRIN, 3-PHOSPHOINOSITIDES, INOSITOL TETRAKISPHOSPHATE 2 SIGNAL TRANSDUCTION PROTEIN, ADAPTOR PROTEIN	SIGNALING PROTEIN ARFI GUANINE NUCLEOTIDE EXCHANGE FACTOR AND PH DOMAIN	SIGNALING PROTEIN ARF! GUANINE NUCLEOTIDE EXCHANGE FACTOR AND PH DOMAIN
Соптроипд	HUMAN SOS 1; CHAIN: A;	BETA-SPECTRIN; IDRO 6 CHAIN: NULL; IDRO 7	HEPATOCYTE GROWTH FACTOR-REGULATED TYROSINE CHAIN: A;	RHO-GEF VAV; CHAIN: A;	DUAL ADAPTOR OF PHOSPHOTYROSINE AND 3- CHAIN: A;	DUAL ADAPTOR OF PHOSPHOTYROSINE AND 3- CHAIN: A;	DUAL ADAPTOR OF PHOSPHOTYROSINE AND 3- CHAIN: A;	DUAL ADAPTOR OF PHOSPHOTYROSINE AND 3- CHAIN: A;	GRP1; CHAIN: A;	GRP1; CHAIN: A;
SeqFold score										
PMF	_	0.21	<u> </u>	1	0.94	0.29	0.99	0.98	0.21	0.98
Verify score	0.12	0.22	0.07	0.14	0.52	4.0-	0.63	0.67	-0.18	0.27
PSI- BLAST	7.80E-58	9.10E-09	9.00E-11	5.40E-24	3.60E-11	1.00E-09	2.60E-18	1.80E-11	9.10E-08	1.10E-15
End	576	820	674	454	816	574	817	816	574	821
Start	263	736	618	260	726	484	719	722	484	725
Chain ID	A		¥	¥	<	4	∢	<	<	. ✓
PDB ID	1dbh	1dro	1dvp	165x	1fao	1168	1fb8	1168	1fgy	1fgy
SEQ B	788	788	788	788	788	788	788	788	788	788

PDB annotation	SIGNALING PROTEIN ARFI GUANINE NUCLEOTIDE EXCHANGE FACTOR AND PH DOMAIN		SIGNAL TRANSDUCTION IRS-1; BETA-SANDWHICH, SIGNAL TRANSDUCTION	TRANSPORT PROTEIN FYVE DOMAIN, ENDOSOME MATURATION, INTRACELLULAR TRAFFICKING, 2 TRANSPORT PROTEIN	COMPLEX (GTP-BINDING/EFFECTOR) RAS-RELATED PROTEIN RAB3A; COMPLEX (GTP-BINDING/EFFECTOR), G PROTEIN, EFFECTOR, RABCDR, 2 SYNAPTIC EXOCYTOSIS, RAB PROTEIN, RAB3A, RABPHILIN	RECEPTOR RECEPTOR, SIGNAL TRANSDUCER OF IL-6 TYPE CYTOKINES, THIRD 2 N-TERMINAL DOMAIN, TRANSMEMBRANE, GLYCOPROTEIN	RECEPTOR RECEPTOR, SIGNAL TRANSDUCER OF IL-6 TYPE CYTOKINES, THIRD 2 N-TERMINAL DOMAIN, TRANSMEMBRANE, GLYCOPROTEIN	RECEPTOR RECEPTOR, SIGNAL TRANSDUCER OF IL-6 TYPE CYTOKUNES, THIRD 2 N-TERMINAL DOMAIN, TRANSMEMBRANE, GLYCOPROTEIN
Coumpound	GRP1; CHAIN: A:	PHOSPHORYLATION PLECKSTRIN (N-TERMINAL PLECKSTRIN HOMOLOGY DOMAIN) MUTANT 1PLS 3 WITH LEU GLU (HIS)6 ADDED TO THE C TERMINUS 1PLS 4 (INS(G105-LEHHHHHH)) (NMR, 25 STRUCTURES) 1PLS 5	INSULIN RECEPTOR SUBSTRATE 1; CHAIN: A, B;	PHOSPHATIDYLINOSITOL-3- PHOSPHATE BINDING FYVE CHAIN: A;	RAB-3A; CHAIN: A; RABPHILIN-3A; CHAIN: B;	GP130; CHAIN: NULL;	GP130; CHAIN: NULL;	GP130; CHAIN: NULL;
SeqFold score								
PMF	90.0	0.93	0.76	0.94	0.1	0.09	0.13	-0.01
Verify score	-0.15	1 .0	0.64	0.04	0.11	-0.08	-0.09	0.14
PSI- BLAST	7.80E-16	1.80E-11	3.60E-05	7.20E-08	3.90E-21	1.10E-08	3.60E-11	7.20E-11
End	814	818	816	671	674	360	477	280
Start AA	.729	728	725	620	290	. 589	387	484
Chain	A		A	<	a			
PDB ID	1fgy	1pls	Iqqg	lvfy	pqz1	16j8	1bj8	15j8
SEQ D	788	788	788	788	788	789	789	789

PDB annotation	CONNECTIN A71, CONNECTIN; TITIN, CONNECTIN, FIBRONECTIN TYPE III	CONNECTIN A71, CONNECTIN; TITIN, CONNECTIN, FIBRONECTIN TYPE III	SIGNALING PROTEIN CYTOKINE RECEPTOR, GLYCOPROTEIN 130, GP130, INTERLEUKINE 6 2 RECEPTOR BETA SUBUNIT, SIGNALING PROTEIN	SIGNALING PROTEIN CYTOKINE RECEPTOR, GLYCOPROTEIN 130, GP130, INTERLEUKINE 6 2 RECEPTOR BETA SUBUNIT, SIGNALING PROTEIN	SIGNALING PROTEIN CYTOKINE RECEPTOR, GLYCOPROTEIN 130, GP130, INTERLEUKINE 6 2 RECEPTOR BETA SUBUNIT, SIGNALING PROTEIN		HORMONE/GROWTH FACTOR/HORMONE RECEPTOR 4- HELICAL BUNDLE, ALPHA HELICAL HONDLE, TERNARY COMPLEX, FN 2 III DOMAINS, BETA SHEET DOMAINS, CYTOKINE-RECEPTOR COMPLEX			
	CONNECT	CONNECT	RECEPTOI GP130, IN BETA SUE	RECEPTO GP130, IN BETA SUE	RECEPTO GP130, IN BETA SUE		HORMON FACTOR/I HELICAL BUNDLE, III DOMAI			
Coumpound	TITIN; CHAIN: NULL;	TITIN; CHAIN: NULL;	GP130; CHAIN: A, B;	GP130; CHAIN: A, B;	GP130; CHAIN: A, B;	NEURAL'ADHESION MOLECULE DROSOPHILA NEUROGLIAN (CHYMOTRYPTIC FRAGMENT CONTAINING THE ICFB 3 TWO AMINO PROXIMAL FIBRONECTIN TYPE III REPEATS ICFB 4 (RESIDUES 610 - 814)) ICFB 5	PLACENTAL LACTOGEN; CHAIN: A; PROLACTIN RECEPTOR; CHAIN: B, C;	CELL ADHESION PROTEIN FIBRONECTIN CELL- ADHESION MODULE TYPE III-10 1FNA 3	CELL ADHESION PROTEIN FIBRONECTIN CELL- ADHESION MODULE TYPE III-10 1FNA 3	CELL ADHESION PROTEIN FIBRONECTIN CELL-
SeqFold score										
PMF score	0.11	0.25	0.06	-0.14	-0.03	0.24	0.19	0.19	0.22	0.41
Verify score	-0.31	-0.02	-0.19	60.0	0.11	0.03	-0.26	-0.14	0.04	0.07
PSI- BLAST	3.60E-07	3.60E-10	7.20E-21	9.00E-15	1.80E-18	1.80E-26	1.60E-14	3.60E-07	1.10E-11	1.30E-12
End AA	360	578	360	490	595	283	359	360	582	989
Start	288	485	195	288	386	385	961	295	492	504
Chain			¥	4	¥		В			
PDB ID	1bpv	1bpv	16qu	1bqu	1 bqu	1cfb	166	lfna	1fna	1fna
SEQ EQ	789	789	789	789	789	789	789	789	789	789

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PDB annotation		CELL ADHESION PROTEIN RGD, EXTRACELLULAR MATRIX IFNF 18	CELL ADHESION PROTEIN KGD, EXTRACELLULAR MATRIX IFNF 18	CELL ADHESION PROTEIN RGD, EXTRACELLULAR MATRIX 1FNF 18	CELL ADHESION PROTEIN RGD, EXTRACELLULAR MATRIX IFNF 18	HEPARIN AND INTEGRIN BINDING HEPARIN AND INTEGRIN BINDING	HEPARIN AND INTEGRIN BINDING HEPARIN AND INTEGRIN BINDING	HYDROLASE TYROSINE PHOSPHATEASE, LAR PROTEIN	HYDROLASE TYROSINE PHOSPHATEASE, LAR PROTEIN	HYDROLASE TYROSINE PHOSPHATEASE, LAR PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN, RGD, EXTRACELLULAR MATRIX, 2 HEPARIN-BINDING, GLYCOPROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN, RGD, EXTRACELLULAR MATRIX, 2 HEPARIN-BINDING, GLYCOPROTEIN	STRUCTURAL PROTEIN IN ECKIN, HEMIDESMOSOME, FIBRONECTIN, CARCINOMA, STRUCTURAL 2 PROTEIN	STRUCTURAL PROTEIN INTEGRIN, HEMIDESMOSOME, FIBRONECTIN, CARCINOMA, STRUCTURAL 2 PROTEIN	STRUCTURAL PROTEIN INTEGRIN, HEMIDESMOSOME, FIBRONECTIN, CARCINOMA, STRUCTURAL 2 PROTEIN	STRUCTURAL PROTEIN TENASCIN,
Coumpound	ADHESION MODULE TYPE III-10 IFNA 3	FIBRONECTIN; 1FNF 6 CHAIN: NULL; 1FNF 7	FIBRONECTIN; 1FNF 6 CHAIN: NULL; 1FNF 7	FIBRONECTIN; IFNF 6 CHAIN: NULL; 1FNF 7	FIBRONECTIN; IFNF 6 CHAIN: NULL; IFNF 7	FIBRONECTIN; CHAIN: A;	FIBRONECTIN; CHAIN: A;	LAR; CHAIN: A, B;	LAR; CHAIN: A, B;	LAR; CHAIN: A, B;	FIBRONECTIN; CHAIN: NULL;	FIBRONECTIN; CHAIN: NULL;	INTEGRIN BETA-4 SUBUNIT; CHAIN: A, B;	INTEGRIN BETA 4 SUBUNIT; CHAIN: A, B;	INTEGRIN BETA-4 SUBUNIT; CHAIN: A, B;	TENASCIN; CHAIN: A, B;
SeqFold score			120.84													
PMF	·	0.51		0.31	-0.07	0.07	0.55	_	-		0.16	0.09	0.1	-1.41	0.92	0.18
Verify		-0.02		-0.03	0.02	0.11	0.05	99.0	0.34	0.72	0.12	-0.13	0.18	0.28	0.3	0.12
PSI- BLAST		5.40E-39	5.40E-39	9.00E-38	9.00E-39	1.10E-19	7.20E-32	0	3.60E-81	0	5.40E-23	1.80E-25	1.10E-17	1.40E-25	2.60E-29	9.00E-17
End		282	588	674	752	473	673	1462	1164	1462	474	582	478	288	584	480
Start		192	194	288	386	196	389	893	782	924	292	388	290	388	390	292
Chain						V	∀	V	В	В			4	∢	4	4
PDB		1fhf	1thf	1fuf	1fbf	Ifish	Ifah	Ilar	Ilar	Ilar	1mfn	1mfn	1983	1983	1983	lara
SEQ	Ö	789	789	789	789	789	789	789	789	789	789	789	789	789	789	789

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PDB annotation	FIBRONECTIN TYPE-III, HEPARIN, EXTRACELLULAR 2 MATRIX, ADHESION, FUSION PROTEIN, STRUCTURAL PROTEIN	STRUCTURAL PROTEIN TENASCIN, FIBRONECTIN TYPE-III, HEPARIN, EXTRACELLULAR 2 MATRIX, ADHESION, FUSION PROTEIN, STRUCTURAL PROTEIN	RECEPTOR DI; RECEPTOR, PHOSPHATASE, SIGNAL TRANSDUCTION, ADHESION, 2 HYDROLASE	RECEPTOR DI; RECEPTOR, PHOSPHATASE, SIGNAL TRANSDUCTION, ADHESION, 2 HYDROLASE				IMMUNE SYSTEM CD32; RECEPTOR, FC, CD32, IMMUNE SYSTEM	PROTEIN BINDING ED-B, FIBRONECTIN, TYPEIII DOMAIN, ANGIOGENESIS, PROTEIN 2 BINDING	PROTEIN BINDING ED-B, FIBRONECTIN, TYPEIII DOMAIN, ANGIOGENESIS, PROTEIN 2 BINDING	TYROSINE PHOSPHATASE SYP, SHPTP-2; TYROSINE PHOSPHATASE, INSULIN SIGNALING, SH2 PROTEIN	
Coumpound		TENASCIN; CHAIN: A, B;	RECEPTOR PROTEIN TYROSINE PHOSPHATASE MU; CHAIN: A, B;	RECEPTOR PROTEIN TYROSINE PHOSPHATASE MU; CHAIN: A, B;	CELL ADHESION PROTEIN TENASCIN (THIRD FIBRONECTIN TYPE III REPEAT) ITEN 3	CELL ADHESION PROTEIN TENASCIN (THIRD FIBRONECTIN TYPE III REPEAT) 1TEN 3	GLYCOPROTEIN FIBRONECTIN (TENTH TYPE III MODULE) (NMR, 36 STRUCTURES) 1TTF 3	FC GAMMA RIIB; CHAIN: A;	FIBRONECTIN; CHAIN: A;	FIBRONECTIN; CHAIN: A;	SHP-2; CHAIN: A, B;	1
SeqFold score			405.81									
PMF score		0.41		-	0.03	0.03	0.31	-0.2	0.98	0.16	-	
Verify score		0.32		0.85	0.08	0.08	0.16	0.03	0.35	0.31	0.44	
PSI- BLAST		1.30E-20	1.40E-89	1.40E-89	1.60E-07	1.30E-08	5.40E-13	1.40E-10	1.60E-05	1.80E-08	1.40E-72	
End		584	1166	1165	584	584	582	163	480	584	1164	
Start AA		390	888	168	487	502	485	27	391	490	835	
Chain		4	٧	٧				4	A	Ą	A	
EDB GI		1qr4	1.pm	lrpm	Iten	1ten	1tt	2fcb	2fnb	2fnb	2shp	
S a S		789	789	789	789	789	789	682	789	682	789	

												
PDB annotation	OXIDOREDUCTASE PDZ DOMAIN, NNOS, NITRIC OXIDE SYNTHASE	OXIDOREDUCTASE PDZ DOMAIN, NNOS, NITRIC OXIDE SYNTHASE	PEPTIDE RECOGNITION PEPTIDE RECOGNITION, PROTEIN LOCALIZATION	PEPTIDE RECOGNITION PEPTIDE RECOGNITION, PROTEIN LOCALIZATION	SIGNAL TRANSDUCTION PROTEIN	SIGNALING PROTEIN DAPPI, PHISH, BAM32; PLECKSTRIN, 3- PHOSPHOINOSITIDES, INOSITOL TETRAKISPHOSPHATE 2 SIGNAL TRANSDUCTION PROTEIN, ADAPTOR	SIGNALLING PROTEIN ARF! GUANINE NUCLEOTIDE EXCHANGE FACTOR AND PH DOMAIN	CYTOKINE LCF, CYTOKINE, LYMPHOCYTE CHEMOATTRACTANT FACTOR, PDZ DOMAIN	OXIDOREDUCTASE BETA-FINGER	OXIDOREDUCTASE BETA-FINGER	MEMBRANE PROTEIN/OXIDOREDUCTASE BETA- FINGER, HETERODIMER	MEMBRANE PROTEIN/OXIDOREDUCTASE BETA- FINGER, HETERODIMER
Coumpound	NEURONAL NITRIC OXIDE SYNTHASE; CHAIN: A; HEPTAPEPTIDE; CHAIN: B;	NEURONAL NITRIC OXIDE SYNTHASE; CHAIN: A; HEPTAPEPTIDE; CHAIN: B;	PSD-95; CHAIN: A; CRIPT; CHAIN: B;	PSD-95; CHAIN: A; CRIPT; CHAIN: B;	BETA-SPECTRIN; 1BTN 4 CHAIN: NULL; 1BTN 5	DUAL ADAPTOR OF PHOSPHOTYROSINE AND 3- CHAIN: A;	GRP1; CHAIN: A;	INTERLEUKIN 16; CHAIN: NULL;	NEURONAL NITRIC OXIDE SYNTHASE (RESIDUES 1- 130); CHAIN: A:	NEÚRONAL NITRIC OXIDE SYNTHASE (RESIDUES 1- 130): CHAIN: A:	ALPHA-I SYNTROPHIN (RESIDUES 77-171); CHAIN: A, NEURONAL NITRIC OXIDE SYNTHASE (RESIDUES I-130); CHAIN: B:	ALPHA-I SYNTROPHIN (RESIDUES 77-171); CHAIN: A; NEURONAL NITRIC OXIDE SYNTHASE
SeqFold score	58.72		57.29								77.08	
PMF		0.23		1	0.04	-0.13	0.05	0.43	0.99	0.94		
Verify score		0.46		1.06	-0.05	0.24	-0.65	0.09	0.5	0.47		0.87
PSI- BLAST	1.40E-11	1.40E-11	9.00E-16	9.00E-16	3.60E-10	5.40E-09	0.0052	3.60E-11	3.60E-10	2.60E-23	3.90E-23	3.90E-23
End	176	171	163	143	388	385	267	138	166	168	140	136
Start	48	52	45	49	299	295	199	27	54	56	54	56
Chain	A	4	A	A		4	V		4	∀	<	«
PDB	1589	1589	1be9	1be9	1btn	1fao	Ifgy	1116	Iqau	1qau	lqav	lqav
SEQ	793	793	793	793	793	793	793	793	793	793	793	793

PDB annotation		MEMBKANE PROTEIN/OXIDOREDUCTASE BETA- FINGER, HETERODIMER	A COT LY CALL CASES	RNA-BINDING PROTEINIRNA I KA PRE-MRNA; SPLICING REGULATION, RNP DOMAIN, RNA COMPLEX	GENE REGULATION/RNA POLY(A)	BINDING PROTEIN 1, PABY 1; NAM, PROTEIN-RNA COMPLEX, GENE	REGULATION/KNA	RNA BINDING PROTEIN RNA- BINDING DOMAIN	NUCLEAR PROTEIN	HETEROGENEOUS NUCLEAR RIBONUCLEOPROTEIN A1, NUCLEAR PROTEIN, HIRNP, RBD, RRM, RNP,	RNA BINDING, 2 RIBONUCLEOPROTEIN	NUCLEAR PROTEIN HETEROGENEOUS NUCLEAR	RIBONUCLEOPROTEIN A1, NUCLEAR	PROTEIN, HUKNY, KBD, KKW, KWY, RNA BINDING, 2 RIBONUCLEOPROTEIN	COMPLEX (RIBONUCLEOPROTEIN/RNA)			1	RNA BINDING PROTEIN RNA-	BINDING DOMAIN
Coumpound	N. B;	ALPHA-1 SYNTROPHIN (RESIDUES 77-171), CHAIN: A; NEURONAL NITRUC OXIDE SYNTHASE (RESIDIES 1-130), CHAIN: B;	Н	SXI-LETHAL PROTEIN, CHAIN: A. B. RNA (5'- R(P*GP*UP*UP*UP*UP*U P*UP*UP*UP*UP*U)- CHAIN: P. O.	<u> </u>		R(*AP*AP*AP*AP*AP*AP*AP *AP*AP*AP*A)-3'); CHAIN:	HU ANTIGEN C. CHAIN: A;	HNRNP A1: CHAIN: NULL:			HNRNP AI; CHAIN: NULL;			U1A SPLICEOSOMAL PROTEIN: 1URN 5 CHAIN: A.	B, C; IURN 6 RNA 21 MER	HAIRPIN (5'-	P* IURN II CHAIN: P, Q, R	MATISASETTI CHAIN: A:	MUSASDII, כחמוזי ה.
SeqFold																				
PMF		-		0.81	0.99			0.71	63.0	3		0.99			0.94				90	 88
Verify score		0.93		0.84	0.41			0.39	0 7 0	ê 5		99.0			0.55				-	0.82
PSI- BLAST		3.60E-16		1.80E-16	5 40E-18			1.30E-15	2000	7.00E-23		1.80E-15			2.60E-15				100	1.60E-15
. End		139		108	100	}		105		<u> </u>		107			112				-	
Start		56		33	•	0		34		-		34			33					35.
Chain		A		4		τ		4							4					۷.
PDB	1	Iqav		167f		<u> </u>		1d8z		Jhal		1ha1			lum					2mss
SEQ	ÖZ	793		794	200	*		794		794		794			794					794

PDB annotation	RNA-BINDING DOMAIN RNA- BINDING DOMAIN, ALTERNATIVE SPLICING	COMPLEX (RIBONUCLEOPROTEIN/DNA) HNRNP A1, UP1; COMPLEX (RIBONUCLEOPROTEIN/DNA), HETEROGENEOUS NUCLEAR 2 RIBONUCLEOPROTEIN A1	RNA BINDING DOMAIN RNA BINDING DOMAIN, RBD, RNA RECOGNITION MOTIF, RRM, 2 SPLICING INHIBITOR, TRANSILATIONAL INHIBITOR, SEX 3 DETERMINATION, X CHROMOSOME DOSAGE COMPENSATION		COMPLEX (TRANSCRIPTION REGULATION/DNA) SREBF-1A; STEROL REGULATORY ELEMENT	BINDING PROTEIN, 2 BASIC-HELIX-	SCHELIA TEANSCRIPTION 3 FACTOR, COMPLEX (TRANSCRIPTION REGILI ATTON/DNA)	CONDITION OF VICTORIAN	REGULATION/DNA) SREBP-1A;	STEROL REGULATORY ELEMENT RINDING PROTEIN 2 BASIC-HELIX-	LOOP-HELIX-LEUCINE ZIPPER,	SREBP, TRANSCRIPTION 3 FACTOR,	COMPLEX (TRANSCRIPTION REGIT ATTON/DNA)	COMPLEX (DNA-BINDING	PROTEIN/DNA) MYN PROTEIN; MAX,	DNA BINDING, BASIC-HELIX-LOOP-	HELIX-LEUCINE ZIPPER, 2	TRANSCRIPTION FACTOR, COMPLEX	COMPLEX (DNA-BINDING PROTEIN/DNA) UPSTREAM
Conmpound	SEX-LETHAL PROTEIN; CHAIN: NULL;	HETEROGENEOUS NUCLEAR RIBONUCLEOPROTEIN A1; CHAIN: A; 12-NUCLEOTIDE SINGLE-STRANDED TELOMETRIC DNA; CHAIN: B;	SEX-LETHAL; CHAIN: A, B, C,		STEROL REGULATORY ELEMENT BINDING PROTEIN 1A; CHAIN: A, B, C,	D; DNA; CHAIN: E, F, G, H;		Vacati naci per	ELEMENT BINDING	PROTEIN 1A; CHAIN: A, B, C,	, 5, 5, 11, 5, 11, 5, 11, 5, 11, 5, 11, 5, 11, 5, 11, 5, 11, 5, 11, 5, 11, 5, 11, 5, 11, 5, 11, 5, 11, 5, 11, 5			MAX PROTEIN; CHAIN: A, C;	DNA; CHAIN: B, D;			•	USF; CHAIN: A, B; DNA; CHAIN: C, D;
SeqFold score							•												
PMF	99.0	0.23	0.96		0.04			200	77.0					-0.03					0.07
Verify score	0.31	0.68	0.61		0.23				† · · ·					0.35					-0.08
PSI- BLAST	1.80E-16	5.40E-30	5.40E-16		2.60E-13	•		1 205 16	1.305-13					9.00E-15					1.80E-13
End	108	=	105		131			9:	611					119					115
Start	33	-	33		65			2	<u>. </u>					57					55
Chain		<	<		⋖			٥	ο					¥					¥
PDB TD	2sxl	2up1	3sxl	_	lam 9		, ,	100	1 9					lan2			_		1an4
SEQ B B SE	794	794	794	_	795			30,	3					795					795

								_
PDB annotation	STIMULATORY FACTOR 1; USF, DNA BINDING, BASIC-HELIX-LOOP-HELIX, LEUCINE ZIPPER, 2 TRANSCRIPTION FACTOR, COMPLEX (DNA-BINDING PROTEIN/DNA)	COMPLEX (TRANSCRIPTION FACTOR MAX/DNA) TRANSCRIPTIONAL REGULATION, DNA BINDING, COMPLEX 2 (TRANSCRIPTION FACTOR MAX/DNA)	COMPLEX (TRANSCRIFTION FACTOR MAX/DNA) TRANSCRIFTIONAL REGULATION, DNA BINDING, COMPLEX 2 (TRANSCRIPTION FACTOR MAX/DNA)		COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) ZINC
Coumpound		TRANSCRIPTION FACTOR MAX; CHAIN: A, B; DNA (5'- D(*CP*AP*CP*CP*GP *TP*GP*GP*T)-3', CHAIN: C, D;	TRANSCRIPTION FACTOR MAX; CHAIN: A, B; DNA (5'- D(*CP*AP*CP*CP*AP*CP*GP *TP*GP*GP*T)-3', CHAIN: C, D;	TRANSCRIPTION ACTIVATIONDNA MYOD BASIC-HELIX-LOOP-HELIX (BHILH) DOMAIN IMDY 3 (RESIDUES 102 - 166) MUTANT WITH CYS 135 REPLACED BY SER IMDY 4 (C135S) COMPLEXED WITH DNA IMDY 5 (5'- D(*TP*CP*AP*CP*AP*GP *CP*TP*GP*TP*TP*GP*A)-3') IMDY 6	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	DNA; CHAIN: A, B, D, E;
SeqFold score						* *		
PMF		0.17	0.05	0.11	-0.19	0.55	0.42	89.0
Verify score		0.06	-0.17	0.15	0.07	0.24	0.31	0.14
PSI- BLAST		3.60E-15	7.20E-15	1.40E-14	1.40E-22	3.60E-26	5.20E-32	5.40E-44
End		119	119	114	405	433	434	433
Start		23	55	53	327	353	357	352
Chain		4	æ	m	V	∢ .	∢	O
PDB D		1hio	1hlo	y y	lalh	lalh	lalh	Ime
SEQ	SON SON SON SON SON SON SON SON SON SON	795	795	795	800	800	800	800

									 -
PDB annotation	FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGERODNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGERUDA) LINC FROER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGERDNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA FINERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA
Coumpound	CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B. D. E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER
SeqFold score									
PMF		-	-	1	1	-	-	1	
Verify score		0.49	0.45	0.52	0.52	0.48	0.45	0.15	61.0
PSI- BLAST		1.80E-46	3.90E-48	2.60E-48	5.40E-47	9.00E-48	1.80E-48	9.00E-50	1.80E-50
End		461	461	489	489	517	545	573	109
Start		380	381	408	408	436	464	492	520
Chain		U	U	ပ	O	O	U	U	ပ
808 E	>	r Ime	Ime y	Ime y	y y	Jme y	Ime y	Jme y	1me
DES B		008	008	008	800	800	800	008	800

PDB annotation	INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGERDINA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGERDNA)	COMPLEX (ZNC FINGERDNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA FINERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA FINTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC PINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2
Coumpound	PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A. B. D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;
SeqFold score				107.23					
PMF		_	1		-	-	-		-
Verify score		0.3	0.21		0.38	0.54	0.54	0.41	0.19
PSI- BLAST		5.40E-51	3.60E-51	3.60E-51	5.40E-51	1.20E-51	1.40E-50	1.40E-50	9.00E-46
End		629	657	658	685	713	713	741	765
Start	-	548	576	576	604	632	632	099	889
Chain .	,	U	υ	ပ	U	U	O	U	U
PDB	+	r 1me	Jme y	Ime y	Jme y	J me	Jme y	1me y	J me
SEQ	ö	800	008	008	800	800	800	800	800

PDB annotation	REGULATION/DNA) YING-YANG 1; TRANSCRETION MITIATOR, MITIATOR ELEMENT, YYL, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; REANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (1 KANDSCKIF ILON REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATIOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DINA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YNG-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)
Coumpound	ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;	YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;	YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;
SeqFold score						
PMF			-	-	0.96	0.93
Verify score		0.05	0.51	0.1	0.12	0.33
PSI- BLAST		1.30E-45	1.30E-57	7.80E-55	1.30E-53	9.00E-35
End		461	489	545	601	629
Start AA		357	385	434	490	528
Chain		O	U	ပ	O	U
PDB		Iubd	1ubd	lubd	1ubd	1ubd
SEQ	Ö	800	800	008	800	800

PDB annotation	COMPLEX (TRANSCRIPTION REGULATIONIDNA) YING-YANG 1; TRANSCRIPTION INTITATION, INTITATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATIONIDNA)	COMPLEX (TRANSCRIPTION REGULATION/DIA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN FECCONITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION RECULATION/DNA) YING-YANG 1; TRANSCRIPTION INITATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN FECCONITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG I; TRANSCRIPTION INITIATION, INITIATION, FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX
Coumpound	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INTIATOR ELEMENT DNA; CHAIN: A, B;	YYI, CHAIN: C, ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI, CHAIN: C; ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B:	YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;
SeqFold score		91.29				
PMF	-		-	66.0	-	
Verify score	0.11		0.22	0.25	0.13	0.15
PSI- BLAST	1.30E-58	5.20E-60	3.60E-35	5.20E-60	1.60E-34	3.90E-61
End	657	828	657	713	713	741
Start	546	548	556	602	612	630
Chain	U	U	U	U	ပ	ပ
PDB CI	lubd	lubd	Iubd	Jubd	lubd	lubd
SEQ S	800	800	800	800	800	800

PDB annotation	(TRANSCRIPTION REGULATION/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	STRUCTURAL PROTEIN TWO REPEATS OF SPECTRIN, ALPHA HELICAL LINKER, REGION, 2.2
Coumpound		ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GL11; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLI1; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLI1; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ALPHA SPECTRIN; CHAIN: A, B, C;
SeqFold			105.23								
PMF		0.93		0.88	0.81	-	6.0	-	1	0.98	0.37
Verify score		0.44		0.18	0.06	0.15	0.16	0.11	0.32	0.41	-0.2
PSI- BLAST		1.30E-43	3.90E-71	2.60E-70	3.90E-71	5.20E-73	1.30E-34	5.20E-79	5.40E-34	1.10E-33	0.0052
End AA		463	519	547	603	631	929	743	712	743	302
Start		367	380	381	408	464	528	576	584	612	16
Chain		4	V.	4	4	∢	∢	V	v V	<	A
PDB ID		2gli	2gli	2gli	2gli	2gli	2gli	2gli	2gli	2gli	lcun
SEQ B B		800	800	800	800	008	800	800	800	800	804

PDB annotation	TANDEM 3-HELIX COILED-COILS, STRUCTURAL PROTEIN	CELL ADHESION FOUR-HELIX BUNDLE	CELL ADHESION FOUR-HELIX BUNDLE	RIBOSOME, HINGE VARIABILITY RIBOSOME, HINGE VARIABILITY	CHAPERONE ARCHAEAL PROTEIN	GROWTH FACTOR (ABU6, 20) MEGF4- 48; GROWTH FACTOR, MURINE EPIDERMAL GROWTH FACTOR, DISULFIDE 2 CONNECTIVITIES, EGF- LIKE DOMAIN, REPEAT	COMPLEX (BLOOD COAGULATION/INHIBITOR) AUTOPROTHROMBIN IIA; HYDROLASE, SERINE PROTEINASE), PLASMA CALCIUM BINDING, 2 GLYCOPROTEIN, COMPLEX (BLOOD COAGULATION/INHIBITOR)	BLOOD COAGULATION BLOOD COAGULATION, EGF, HYDROLASE, SERNE PROTEASE	MEMBRANE PROTEIN LECTIN-LIKE, NEUROBIOLOGY, CELL-CELL ADHESION, CELL-CELL 2 RECOGNITION, ALTERNATIVE SPLICING, MEMBRANE PROTEIN	TRANSPORT PROTEIN SHBG; STEROID TRANSPORT, LAMININ G- LIKE DOMAIN, JELLYROLL, 2 ANDROGEN BINDING PROTEIN (ABP), SEX STEROID BINDING PROTEIN 3 (SBP)	HYDROLASE/HYDROLASE INHIBITOR PROTEIN-PEPTIDE COMPLEX
Coumpound		ALPHA-CATENIN; CHAIN: A;	ALPHA-CATENIN; CHAIN: A; BETA-CATENIN; CHAIN: B;	RIBOSOME RECYCLING FACTOR; CHAIN: A;	PREFOLDIN; CHAIN: A; PREFOLDIN; CHAIN: B; PREFOLDIN; CHAIN: C;	EPIDERMAL GROWTH FACTOR; CHAIN: NULL;	ACTIVATED PROTEIN C; CHAIN: C, L; D-PHE-PRO- MAI; CHAIN: P;	FACTOR VII; CHAIN: NULL;	NEUREXIN-J BETA; CHAIN: A, B, C. D. E. F. G, H;	SEX HORMONE-BINDING GLOBULIN; CHAIN: A;	DES-GLA FACTOR VIIA (HEAVY CHAIN); CHAIN: H, I; DES-GLA FACTOR VIIA
SeqFold score											
PMF		-	1	90.0	0.01	0.58	-0.12	0.99	9.4	0.72	0.05
Verify		0.41	0.37	-0.65	-0.34	0.39	0.55	96.0	0.26	0.46	0.48
PSI- BLAST		3.90E-59	6.50E-70	0.0072	1.30E-06	6.50E-12	6.50E-19	1.30E-11	1.30E-26	3934 2.60E-27	3.90E-21
End		257	257	441	103	3984	4033	3987	3928	3934	4033
Start		79	54	367	S	3949	3943	3949	3771	3773	3944
Chain		V	4	A	V V		٦.		₹	٠ ح	1
PDB		Idov	9 ₂	1eh1	15¢k	la3p	laut	1bf9	1c4r	1d2s	ldva
SEQ.	Ö	804	804	804	804	808	808	808	808	808	808

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PDB annotation		HYDROLASEHYDROLASE INHIBITOR PROTEIN-PEPTIDE COMPLEX	SERINE PROTEINASE COAGULATION FACTOR II; COAGULATION FACTOR II; FETOMODULIN, TM, CD141 ANTIGEN; EGR-LIKE DOMAINS, PROTEINASE, EGF-LIKE DOMAINS, ANTIFIBRINOLYTIC COMPLEX	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCTUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I
Coumpound	(LIGHT CHAIN); CHAIN: L, M; (DPN)-PHE-ARG; CHAIN: C, D; PEPTIDE E-76; CHAIN: X, Y;	DES-GLA FACTOR VIIA (HEAVY CHAIN); CHAIN: H, I, DES-GLA FACTOR VIIA (LIGHT CHAIN); CHAIN: L, M; OPPY-PHE-ARG; CHAIN: C, D; PEPTIDE E-76; CHAIN: X, Y;	THROMBIN LIGHT CHAIN; CHAIN: A, B, C, D; THROMBIN HEAVY CHAIN; CHAIN: M, N, O, P; THROMBOMODULIN; CHAIN: I, J, K, L; THROMBIN INHIBITOR L-GLU-L-GLY-L-ARM; CHAIN: E, F, G, H;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;
SeqFold score								
PMF		-0.18	-0.19	-	1202.08	6.0	0.82	-1.41
Verify score		0.25	90.00	0.26	0.22	0.45	0.27	0.1
PSI- BLAST		1.30E-13	1.30£-11	3.60E-33	1.30E-32	3.60E-21	1.60E-49	1.30E-32
End		4344	4336	1234	1338	1440	1547	1652
Start		4263	4232	1066	1171	1279	1352	1460
Chain ID		1		<	∢	4	∢	4
PDB		ldva	ldx5	ledh	1edh	ledh	ledh	1edh
SEQ S D S		808	808	808	808	808	808	808

PDB annotation	AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12; CADHERIN, CELL ANDESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12: CADHERIN, CELL
Coumpound		E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B:	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;			
SeqFold									
PMF		0.96	0.99	1	0	0.96	0.51	0.78	-
Verify score		0.21	0.19	0.32	0	0.26	0.3	0.24	0.18
PSI- BLAST		3.60E-33	1.60E-28	3.60E-26	1.60E-20	3.60E-29	1.10E-50	1.30E-28	3.60E-48
End		1750	1860	1960	354	2062	2163	2264	2371
Start		1589	0691	1800	. 182	1898	1975	2104	2178
Chain		∢	∢	V		₹	4	<	4
PDB ID		1edh	ledh	1ed h	ledh	ledh	1edh	ledh	1edh
SEQ	ğ	808	808	808	808	808	808	808	808

PDB annotation	ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELLAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECADI2, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM
Coumpound		E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;						
SeqFold score						120.62			
PMF		-	0.86	0.93	96.0		-		0.94
Verify score		0.32	0.07	0	0.1		0.47	0.27	0.31
PSI- BLAST		1.60E-35	1.805-29	1.80E-38	5.40E-32	1.80E-57	1.80E-57	7.20E-35	5.40E-29
End		2473	2577	2683	2789	2895	2898	3003	3105
Start		2306	2414	2488	2619	2692	2693	2831	2941
Chain ID		4	<	<	4	∢	∢	4	¥
PDB		1edh	ledh	ledh	ledh	ledh	1edh	ledh	1edh
SEQ	Ž	808	808	808	808	808	808	808	808

PDB annotation	BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELLAL CADHERIN DOMAINS I AND 2 ECADI 2: CADHERIN, CELL	ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN	EPITHELIAL CADHERIN DOMAINS 1	AND 2, ECAD12; CADRENIN, CELE	BINDING PROTEIN	CELL ADHESION PROTEIN	EPITHELIAL CADHERIN DOMAINS I	ADHESION PROTEIN, CALCIUM	BINDING PROTEIN	CELL ADHESION PROTEIN	AND 2. ECADI2: CADHERIN. CELL	ADHESION PROTEIN, CALCIUM	BINDING PROTEIN	CELL ADHESION PROTEIN	EPITHELIAL CADHEIUN DOMAINS I	AND 2, ECADI2; CADHERIN, CELL	ADHESION PROTEIN, CALCIUM	OF LANDERON DECTEN	CELL ADRESION FRO LEIN	AND 2 ECADIS: CADHERIN. CELL	ADHESION PROTEIN, CALCIUM	BINDING PROTEIN	CELL ADHESION PROTEIN EPITTE: 141 CADHER IN DOMAINS 1	STREET, CANTERN CELL	AND 2, ECAD-12, CADREMIN, CELE	BINDING PROTEIN	CELL ADHESION PROTEIN	EPITHELIAL CADHERIN DOMAINS 1	AND 2, ECAD12; CADHERIN, CELL	ADHESION PROTEIN, CALCIUM BINDING PROTEIN
Coumpound		E-CADHERIN; CHAIN: A, B;		E-CADHERIN; CHAIN: A, B;		-		E-CADHERIN; CHAIN: A, B;				E-CADHERIN; CHAIN: A, B;				E-CADHERIN; CHAIN: A, B;				To the state of th	E-CADHEKIN; CHAIN: A, B;				E-CADHERIN; CHAIN: A, B;				E-CADHERIN; CHAIN: A, B;			
SeqFold																																
PMF	-	0.53		0.64				-1.41				96.0				_					0.3				0.83				0.82			
Verify score		0.38		0.16	2			0.57				0.45				99.0					0.12				0.29				0.17			
PSI- BLAST		7.20E-25		\$ 40F-28	201.0			1.80E-32				1.80E-48				3.60E-30					1.80E-51				1.10E-29				3.60E-29			
End		450		3107) } }			3313				3418				3523					248				556		_	_	662			
Start AA		296		3046	2			3120				3225				3355					39				406				464	<u>.</u>		
Chain		<			ς .			4				A		_	-	4					∢				A					· ·		
PDB	\dagger	ledh		1997	ıını 1			1edi				ledh				ledh					흄				ledh				ledh			
SEQ	ÿ	808		80	90			808	}			808				808					808				808			_	808	}		

PDB annotation	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN			MATRIX PROTEIN EXTRACELLULAR MATRIX, CALCIUM-BINDING, GLYCOPROTEIN, 2 REPEAT, SIGNAL, MULTIGENE FAMILY, DISEASE MUTATION, 3 EGF-LIKE DOMAIN, HUMAN FIBRILLIN-1 FRAGMENT, MATRIX PROTEIN		GLYCOPROTEIN GLYCOPROTEIN	GLYCOPROTEIN GLYCOPROTEIN	CELL ADHESION PROTEIN CADHERIN	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN
Coumpound	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	FIBRILLIN; CHAIN: NULL;	P-SELECTIN; CHAIN: NULL;	LAMININ; CHAIN: NULL;	LAMININ; CHAIN: NULL;	N-CADHERIN; INCG 3	N-CADHERIN; 1NCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3
SeqFold score												
PMF	0.27	-		0.69	66.0	0.88	0.27	-0.03	0.29	69.0	0.29	0.1
Verify score	-0.05	0.33	0.31	0.11	0.71	1.03	-0.1	0.07	0.12	0.13	10:0	0.46
PSI- BLAST	5.40E-22	1.10E-55	1.10E-32	9.00E-32	1.805-16	1.30E-11	1.10E-13	5.40E-21	9.00E-06	0.00014	1.40E-14	0.00018
End	812	917	1022	1129	4022	3988	4050	4075	1127	1232	1439	1546
Start	591	718	854	656	3946	3949	3924	3954	1062	1167	1350	1480
Chain ID	∀	<	<	<								
PDB	1edh	ledh	ledh	ledh	n n	1£b	1klo	1klo	Incg	lncg	Incg	lncg
SEQ	808 808	808	808	808	808	808	808	808	808	808	808	808

PDB annotation	INCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCI 13	CELL ADHESION PROTEIN CADHERIN INCI 13	CELL ADHESION PROTEIN CADHERIN INCI 13
Coumpound		N-CADHERIN; INCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3	N-CADHERIN, INCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3
SeqFold score																					
PMF score		0.09	0.09	0.31	0.1	0.11	0.1	0.74	0.43	0.57	0.7	0.39	0.51	0.34	0.36	8.0	0.53	0.64	0.78	96.0	0.22
Verify score		0.15	-0.13	0.35	0.22	0.28	-0.08	0.41	-0.27	0.34	0.44	0.35	0.45	90'0	0.33	0.16	-0.19	0.38	90:00	90.0	0.43
PSI- BLAST		3.60E-05	9.00E-06	3.60E-06	3.60E-17	0.00036	3.60E-12	1.80E-06	0.00036	1.60E-05	1.80E-19	0.00054	3.60E-06	1.80E-05	1.60E-11	7.20E-20	0.00018	5.40E-05	1.80E-06	5.40E-05	1.30E-13
End		229	1650	1748	2061	2161	2263	2370	2458	2681	2788	2988	3106	3191	3311	811	006	1003	1129	1234	1440
Start AA		155	1599	1667	1970	2079	2178	2304	2411	2593	2692	2913	3039	3120	3225	716	852	932	1065	1172	1350
Chain 13																			В	В	В
PDB ID		lncg	lncg	Incg	lncg	Incg	lncg	lncg	Incg	Incg	Incg	Incg	lncg	1ncg	Incg	Incg	Incg	lncg	lnci	1nci	1nci
Se Se		808	808	808	808	808	808	808	808	808	808	808	808	808	808	808	808	808	808	808	808

PDB annotation	CELL ADHESION PROTEIN CADHERIN INCI 13	CELL ADHESION PROTEIN CADHERIN INCI 13	CELL ADHESION PROTEIN CADHERIN INCI 13	CELL ADHESION PROTEIN CADHERIN INCI 13	CELL ADHESION PROTEIN CADHERIN INCI 13	CELL ADHESION PROTEIN CADHERIN INCI 13	CELL ADHESION PROTEIN CADHERIN INCI 13	CELL ADHESION PROTEIN CADHERIN INCI 13	CELL ADHESION PROTEIN CADHERIN INCI 13	CELL ADHESION PROTEIN CADHERIN INCI 13	CELL ADHESION PROTEIN CADHERIN INCI 13	CELL ADHESION PROTEIN CADHERIN INCI 13	CELL ADHESION PROTEIN CADHERIN INCI 13	CELL ADHESION PROTEIN CADHERIN INCI 13	CELL ADHESION PROTEIN CADHERIN INCI 13	CELL ADHESION PROTEIN CADHERIN INCI 13	CELL ADHESION PROTEIN CADHERIN INCI 13	CELL ADHESION PROTEIN CADHERIN INCI 13	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL
Conmpound	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;
SeqFold score																					
PMF	0.65	9.0	0.4	0.23	0.05	0.1	0.75	0.58	90.0	<u>-</u>	0.92	0.42	0.17	0.49	0.99	96.0	0.71	0.63	_	_	0.54
Verify score	-0.08	0.35	-0.17	0.08	-0.25	-0.11	0.17	-0.14	0	0.56	0.42	0.56	-0.2	0.67	0.87	0.15	-0.4 4	0.36	0.45	0.34	0.3
PSI- BLAST	0.00018	9.00E-07	1.60E-05	1.80E-16	0.0013	1.30E-11	5.40E-07	0.0000	1.60E-05	3.60E-19	0.0036	5.40E-06	0.00036	1.80E-10	1.80E-08	1.80E-19	5.40E-05	1.80E-05	3.60E-36	5.40E-33	1.80E-22
End	1547	1750	248	2062	2163	2264	2371	2458	2683	2789	3003	3105	3191	3313	3418	812	917	1022	1234	1338	1440
Start AA	1491	1667	181	1970	2116	2178	2307	2414	2620	2692	2942	3044	3146	3225	3354	715	862	932	1039	1147	1270
Chain ID	В	В	В	m	В	В	æ	æ	В	В	В	В	В	В	B	В	В	æ	<	4	Ą
PDB UD	Inci	Inci	Inci	Inci	Inci	Inci	Inci	Inci	lnci	Inci	Inci	Inci	Inci	Inci	Inci	Inci	lnci	1nci	Incj	Incj	Inci
SEQ PO	808	808	808	808	808	808	808	808	808	808	808	808	808	808	808	808	808	808	808	808	808

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PDB annotation	ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
Coumpound		N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERM; CHAM: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A:	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;
SeqFold																	120.25				
PMF score		0.99	-	0.21	-	0.74	96.0	0.76	0.33	98.0	_		0.75	0.83	0.3	1			0.99	_	0.58
Verify score		0.29	0.03	0.05	0.45	-0.05	0.31	0.42	0.11	0.48	0.17	0.27	-0.1	0.14	0.25	0.29		0.28	0.31	0.62	-0.03
PSI- BLAST		3.60E-53	1.80E-34	5.40E-24	3.60E-33	3.60E-32	7.20E-27	3.60E-28	7.20E-55	3.60E-30	5.40E-52	5.40E-36	7.20E-32	1.60E-41	3.60E-27	1.30E-32	5.40E-63	5.40E-63	3.60E-38	1.80E-29	1.40E-31
End		1547	1652	354	1750	1981	1960	2062	2163	2264	2371	2473	2577	2683	450	2789	2897	2898	3003	3105	3196
Start		1351	1458	155	1562	1667	1782	1898	1970	2079	2178	2300	2407	2488	256	2593	2691	2693	2825	2913	3039
Chain D		A	A	V	A	4	<	, A	¥	¥	4	A	∀	¥	. V	V	¥	4	¥	¥	¥
PDB DD	1	Incj	lncj	lncj	lncj	Incj	Incj	lncj	lncj	Incj	Incj	Incj	Incj	Incj							
SEQ	ö	808	808	808	808	808	808	808	808	808	808	808	808	808	808	808	808	808	808	808	808

PDB annotation	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	COMPLEX (BLOOD COAGULATION/INHIBITOR) CHRISTMAS FACTOR; COMPLEX,	INHIBITOR, HEMOPHILIANGGH, BLOOD COAGULATION, 2 PLASMA, SERINE PROTEASE, CALCIUM- BINDING, HYDROLASE, 3	GLYCOPROTEIN SEPINE PROTEASE EVILA:	SERINE FRO LEASE FYILY, FYILY, BLOOD COAGULATION, SERINE PROTEASE	SERINE PROTEASE FVIIA; BLOOD COAGULATION, SERINE PROTEASE
Coumpound	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	FACTOR IXA; CHAIN: C, L.; D-PHE-PRO-ARG; CHAIN: I;		COACH ATION EACTOR	COAGULATION FACTOR VIIA (LIGHT CHAIN); CHAIN: L; COAGULATION FACTOR VIIA (HEAVY CHAIN); CHAIN: H; TRIPEPTIDYL INHIBITOR: CHAIN;	COAGULATION FACTOR VIIA (LIGHT CHAIN); CHAIN: L; COAGULATION FACTOR VIIA (HEAVY CHAIN); CHAIN: H; TRIPEPTIDYL
SeqFold score																
PMF score	0.63	0.99	_	0.39	0.21	0.35	0.99	0.18	1	0.88	-	-0.19		9	-0.19	0.57
Verify	0.36	0.48	0.63	0.52	-0.28	0.3	0.37	0.02	0.23	0.34	0.39	0.01		c	7.0	0.68
PSI- BLAST	1.80E-34	7.20E-51	3.60E-32	3.60E-13	1.80E-57	1.80E-34	9.00E-30	1.80E-25	7.20E-62	1.80E-34	1.10E-34	1.40E-10		1 100	11-305-1	1.20E-21
End	3313	3418	3523	3621	248	556	299	812	917	1022	1129	4312		,00,	400	4033
Start	3120	3225	3346	3433	39	390	467	57.1	717	827	932	4224	***	555	3943	3951
Chain	A	<	¥	4	V	V	¥	4	V	4	4	L)		,	ــ	٦
PDB	lncj	Incj	lncj	lncj	1pfx		-	- Тей	lqfk							
SEQ D	808	808	808	808	808	808	808	808	808	808	808	808		8	80.80 80.80	808

PDB annotation	1 11 11 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	SERINE PROTEASE FYILA; FYILA; BLOOD COAGULATION, SERINE PROTEASE			PROTEASE	COAGULATION FACTOR SERINE	COAGULATION, COAGULATION FACTOR	CELL ADHESION UVOMORULIN;	ADHESION	CELL ADHESION UVOMORULIN;	ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
Coumpound	INHIBITOR; CHAIN: C;	COAGULATION FACTOR VIA (LIGHT CHAIN); CHAIN: L; COAGULATION FACTOR	CHAIN: H; TRIPEPTIDYL INHIBITOR; CHAIN: C;	COAGULATION FACTOR VIIA (LIGHT CHAIN); CHAIN:	L; COAGULATION FACTOR VIIA (HEAVY CHAIN); CHAIN: H; TRIPEPTIDYL NURTTOR: CHAIN: C:	COAGULATION FACTOR IX;	CHAIN: A; COAGULATION FACTOR IX; CHAIN: B;	EPITHELIAL CADHERIN;	CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL:	(500)	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;
SeqFold score																	
PMF		-0.17		-0.17		-0.19		0.27		0.93		0.45	0.62	0.13	0.04	0.63	0.07
Verify		0		0.37		0.03		0.24		-0.06		0.04	0.54	0.28	0.37	0.45	0.35
PSI- BLAST		5.40E-14		7.20E-13	. •	7.80E-14		7.80E-20		1.60E-07	•	1.30E-17	1.30E-10	0.0013	1.80E-19	1.30E-08	2.60E-07
End	1	4202		4344		4001		1133		1133		1238	1334	1342	1444	1549	226
Start AA		4121		4267		3951		1041		1066		1145	1249	1279	1350	1455	155
Chain		1		1		В											
PDB TD	1	1 q\$		1qfk		17		Isuh		Isuh		lsuh	1suh	1suh	Isuh	1suh	1suh
SEQ	ö	808		808		808	}	808		808		808	808	808	808	808	808

CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL
EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN: CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;
0.31	0.13	99.0	0.75	0.48	0.45	0.13	0.09	0.4	0.77	0.94	0.58	0.25	0.15
0.41	-0.2	0.07	0.17	-0.06	-0.18	-0.41	0.42	-0.37	0.44	0.35	0.03	-0.04	-0.37
1.00E-12	9.00E-09	1.30E-12	3.60E-06	1.30E-14	3.60E-05	1.80E-06	2.60E-15	1.30E-15	3.90E-20	3.60E-09	3.90E-05	3.60E-05	1.80E-12
1650	1656	1754	1754	1867	1868	252	2163	2268	2375	2375	2475	2477	2581
1570	1589	1667	1690	1771	1800	182	2084	2178	2280	2306	2395	2414	2488
1suh	1suh	1suh	Isuh	1suh	1suh	Ísuh	1suh	1suh	Isuh	1suh	1suh	1suh	1suh
Ö2 808	808	808	808	808	808	808	808	808	808	808	808	808	808
	1suh 1570 1650 1.00E-12 0.41 0.31 EPITHELIAL CADHERIN; CHAIN: NULL;	Isuh 1570 1650 1.00E-12 0.41 0.31 EPITHELIAL CADHERIN; Isuh 1589 1656 9.00E-09 -0.2 0.13 EPITHELIAL CADHERIN; CHAIN: NULL; CHAIN: NULL; CHAIN: NULL; CHAIN: NULL;	1suh 1570 1650 1.00E-12 0.41 0.31 EPITHELIAL CADHERIN; 1suh 1589 1656 9.00E-09 -0.2 0.13 EPITHELIAL CADHERIN; 1suh 1667 1754 1.30E-12 0.07 0.68 EPITHELIAL CADHERIN; CHAIN: NULL; CHAIN: NULL; CHAIN: NULL; CHAIN: NULL;	Isuh 1570 1650 1.00E-12 0.41 0.31 EPITHELIAL CADHERIN; Isuh 1589 1656 9.00E-09 -0.2 0.13 EPITHELIAL CADHERIN; Isuh 1667 1754 1.30E-12 0.07 0.68 EPITHELIAL CADHERIN; Isuh 1690 1754 3.60E-06 0.17 0.75 EPITHELIAL CADHERIN; CHAIN: NULL; CHAIN: NULL; CHAIN: NULL; CHAIN: NULL;	1suh 1570 1656 1.00E-12 0.41 0.31 EPITHELIAL CADHERIN; CHAIN: NULL; 1suh 1589 1656 9.00E-09 -0.2 0.13 EPITHELIAL CADHERIN; CHAIN: NULL; 1suh 1667 1754 1.30E-12 0.07 0.68 EPITHELIAL CADHERIN; CHAIN: NULL; 1suh 1690 1754 3.60E-06 0.17 0.75 CHAIN: NULL; CADHERIN; CHAIN: NULL; 1suh 1777 1867 1.30E-14 -0.06 0.48 EPITHELIAL CADHERIN; CHAIN: NULL;	Isuh 1570 1656 1.00E-12 0.41 0.31 EPITHELIAL CADHERIN; CHAIN: NULL; CHAIN: NULL; CHAIN: NULL; CHAIN: NULL; Isuh 1667 1754 1.30E-12 0.07 0.68 EPITHELIAL CADHERIN; CHAIN: NULL; CHAIN: NULL; CHAIN: NULL; CHAIN: NULL; CHAIN: NULL; CHAIN: NULL; CHAIN: NULL; CHAIN: NULL; CHAIN: NULL; CHAIN: NULL; CHAIN: NULL; CHAIN: NULL; CHAIN: NULL; CHAIN: NULL; CHAIN: NULL; CHAIN: NULL; CHAIN: NULL; CHAIN: NULL; CHAIN: NULL;	15th 1570 1650 1.00E-12 0.41 0.31 EPITHELIAL CADHERIN; CHAIN: NULL; CHAIN: NUL	1suh 1570 1656 1.00E-12 0.41 0.31 EPITHELIAL CADHERIN; CHAIN: NULL; CHAIN: NUL	15uh 1589 1656 9.00E-09 -0.2 0.13 EPITHELIAL CADHERIN; CHAIN: NULL; CHAIN: NUL	15th 1570 1650 1.00E-12 0.41 0.31 EPITHELIAL CADHERIN; CHAIN: NULL; CHAIN: NUL	15th 1570 1650 1.00E-12 0.41 0.31 EPITHELIAL CADHERIN; Contains 1589 1656 9.00E-09 -0.2 0.13 EPITHELIAL CADHERIN; Contains 1667 1754 1.30E-12 0.07 0.68 EPITHELIAL CADHERIN; Contains 15th 1777 1867 1.30E-14 -0.06 0.48 EPITHELIAL CADHERIN; Contains 15th 1777 1867 1.30E-14 -0.06 0.48 EPITHELIAL CADHERIN; Contains 15th	15th 1570 1650 1.00E-12 0.41 0.31 EPITHELIAL CADHERIN; O CHAIN: NULL; O C	18th 1570 1650 1.00E-12 0.41 0.31 EPITHELIAL CADHERIN; O CHAIN: NULL; O C

SEQ	PDB	Chain ID	Start AA	End	PSI- BLAST	Verify	PMF	SeqFold	Coumpound	PDB annotation
SO.										ADHESION
808	Isuh		2489	2581	1.30E-13	91.0	0.98		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UYOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	1suh		2591	2681	2.60E-10	0.02	0.12		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	1suh		2692	2793	3.60E-23	0.42	66.0		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	1suh		2806	2902	2.60E-21	0.43	0.87		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	1 suh		2831	2902	3.60E-09	0.01	0.25		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMOKULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	1suh		2914	3007	1.20E-14	0.37	0.93		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCTUM BINDING, CELL ADHESION
808	lsuh		2941	3007	1.80E-06	-0.07	0.86		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCTUM BINDING, CELL ADHESION
808	1suh		3026	3109	2.60E-21	0.36	0.89		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	1sth		3046	3109	3.60E-06	0.4	0.72		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	Isuh		3120	3185	0.0013	0.1	0.28		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UYOMOKULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	Isuh		3120	3213	3.90E-12	0.43	0.25		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	1suh		3225	3317	1.10E-14	0.6	0.82		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	1suh		3330	3422	3.90E-21	0.58	0.98		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN. CALCIUM BINDING, CELL ADHESION
808	Isuh		3355	3422	1.30E-09	99.0	0.93	R	EPITHELIAL CADHERIN;	CELL ADHESION UVOMORULIN;

PDB Chain Start End D D AA AA	Start		End		PSI- BLAST	Verify score	PMF score	SeqFold score	Coumpound CHAIN: NUIT:	PDB annotation CADHERIN, CALCIUM BINDING, CELL
1suh 3435 3524 6.50E-11	3524	3524	-	6.50E-11		0.56	0.65		CHAIN: NOLL; EPITHELIAL CADHERIN; CHAIN: NULL;	ADHESION CELL ADHESION UVOMORULIN; CELL ADHESION UVOMORULIN; ADHERION
1suh 363 454 0.0001	454 0.0001	454 0.0001	0.0001		1	0.08	0.24		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
1suh 406 454 1.60E-05	454 1.60E-05	454 1.60E-05	1.60E-05		17	9.0	0.19		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
1suh 467 560 1.30E-15 -C	560 1.30E-15	560 1.30E-15	1.30E-15		١٢	-0.03	0.19		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
1suh 590 658 3.90E-05 0.39	658 3.90E-05	658 3.90E-05	3.90E-05		Ö	68	0.57		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
15uh 716 816 1.40E-23 -0.21	816 1.40E-23	816 1.40E-23	1.40E-23		Ģ	= .	0.95		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
1suh 826 921 1.00E-17 0.4	921 1.00E-17	921 1.00E-17	1.00E-17		0.4		0.72		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
921	921 5.40E-07	921 5.40E-07	5.40E-07		0.17	,	0.45		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
930 1026 5.20E-13	1026 5.20E-13	1026 5.20E-13	5.20E-13		0.37		0.57		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
3949 4026	4026 1.30E-18	4026 1.30E-18	1.30E-18		0.2	_	0.59		T-PLASMINOGEN ACTIVATOR F1-G; 1TPG 7 CHAIN: NULL; 1TPG 8	PLASMINOGEN ACTIVATION
1xka L 4267 4348 5.40E-12 0.15	4267 4348 5.40E-12	4348 5.40E-12	5.40E-12		0.15	_	-0.19		BLOOD COAGULATION FACTOR XA; CHAIN: L, C;	BLOOD COAGULATION FACTOR STUART FACTOR, BLOOD COAGULATION FACTOR, SERINE PROTEINASE, EPIDERMAL 2 GROWTH FACTOR LIKE DOMAIN
9wga A 4166 4337 3.60E-10 0.05	4166 4337 3.60E-10	4337 3.60E-10	3.60E-10		0.0	5	-0.2		LECTIN (AGGLUTININ) WHEAT GERM AGGLUTININ (ISOLECTIN 2) 9WGA 3	
laut L 3857 3947 6.50E-19 0.55	3947 6.50E-19	3947 6.50E-19	6.50E-19	П	0	25	-0.12		ACTIVATED PROTEIN C;	COMPLEX (BLOOD

PDB annotation	COAGULATION/NHIBITOR) AUTOPROTHROMBIN IIA; HYDROLASE, SERINE PROTEINASE), PLASMA CALCTUM BINDING, 2 GLYCOPROTEIN, COMPLEX (BLOOD COAGULATION/NHIBITOR)	BLOOD COAGULATION, SERINE PROTEASE, COMPLEX, CO-PACTOR, 2 RECEPTOR ENZYME, INHIBITOR, GLA, EGF, 3 COMPLEX (SERINE PROTEASE/COFACTOR/LIGAND)	BLOOD COAGULATION, SERINE PROTEASE, COMPLEX, CO-PACTOR, 2 RECEPTOR ENZYME, INHIBITOR, GLA, EGF, 3 COMPLEX (SERINE PROTEASE/COFACTOR/LIGAND)	HYDROLASE/HYDROLASE INHIBITOR PROTEIN-PEPTIDE COMPLEX	HYDROLASE/HYDROLASE INHIBITOR PROTEIN-PEPTIDE COMPLEX	HYDROLASE/HYDROLASE INHIBITOR PROTEIN-PEPTIDE COMPLEX
Coumpound	CHAIN: C, L; D-PHE-PRO- MAI; CHAIN: P;	BLOOD COAGULATION FACTOR VIIA; CHAIN: L, H; SOLUBLE TISSUE FACTOR; CHAIN: T, U; D-PHE-PHE- ACG- CHLOROMETHYLKETONE (DFFRCMK) WITH CHAIN: C;	BLOOD COAGULATION FACTOR VILA; CHAIN: I, H; SOLUBLE TISSUE FACTOR; CHAIN: T, U; D-PHE-PHE- ARG- CHLOROMETHYLKETONE (DFFRCMK) WITH CHAIN: C;	DES-GLA FACTOR VIIA (HEAVY CHAIN); CHAIN: H, I; DES-GLA FACTOR VIIA (LIGHT CHAIN); CHAIN: L, M; (DPN)-PHE-ARG; CHAIN: C, D; PEPTIDE E-76; CHAIN: X, Y;	DES-GLA FACTOR VIIA (HEAVY CHAIN); CHAIN: H, I; DES-GLA FACTOR VIIA (LIGHT CHAIN); CHAIN: L, M; (DPN)-PHE-ARG; CHAIN: C, D; PEPTIDE E-76; CHAIN: X, Y;	DES-GLA FACTOR VIIA (HEAVY CHAIN); CHAIN: H, I; DES-GLA FACTOR VIIA (LIGHT CHAIN); CHAIN: L, M; (DPN)-PHE-ARG; CHAIN: C, D; PEPTIDE E-76; CHAIN: X, Y;
SeqFold score						
PMF		0.07	0.07	0.15	0.05	-0.07
Verify score		60:0		-0.08	0.48	0.12
PSI- BLAST		1.40E-12	3.60E-14	1.40E-12	3.90E-21	3.60E-14
End		3897	3976	3897	3947	3976
Start AA		3831	3900	3831	3858	3900
Chain ID			1	1	a a	ı
PDB ID		1dan	1dan	ldva	Idva	1dva
SEQ NO:		808	808	808	808	808

	 _				r			,
PDB annotation	HYDROLASEHYDROLASE INHIBITOR PROTEIN-PEPTIDE COMPLEX	HYDROLASE/HYDROLASE INHIBITOR PROTEIN-PEPTIDE COMPLEX	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECADIS; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELLAL CADHERIN DOMAINS 1 AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I
Coumpound	DES-GLA FACTOR VIIA (HEAVY CHAIN); CHAIN: H, I; DES-GLA FACTOR VIIA (LIGHT CHAIN); CHAIN: L, M; OPN)-PHE-ARG; CHAIN: C, D; PEPTIDE E-76; CHAIN:	DES-GLA FACTOR VIIA (HEAVY CHAIN); CHAIN: H, I; DES-GLA FACTOR VIIA (LIGHT CHAIN); CHAIN: L, M; (DPN)-PHE-ARG; CHAIN: C, D; PEPTIDE E-76; CHAIN: X, Y;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;
SeqFold								
PMF	-0.15	-0.19		96:0	0.93	0.89	0.65	-1.41
Verify score	0.03	0.25	0.35	0.16	0.23	0.3	0.03	0.19
PSI- BLAST	3.60E-12	1.30E-13	1.10E-33	9.00E-30	1.60E-20	3.60E-54	7.20E-32	3.60E-33
End	4168	4259	1234	1338	1440	1547	1652	1750
Start AA	4078	4170	1026	1171	1279	1352	1455	1589
Chain	در	در	∢	4	∢	∢	∀ ·	4
PDB	ldva	ldva	1edh	ledh	1edh	1edh	1edh	ledh
SEQ NO ED	808	808	808	808	808	808	808	808

<u>_</u>	·		<u> </u>						
PDB annotation	AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELLAL CADHERIN DOMAINS I AND 2, ECAD 12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELLAL CADHERIN DOMAINS 1 AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECADIS; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL
Coumpound		E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;
SeqFold score									
PMF score		0.25	0.99	0.07	0.94	0.34	0.92	_	-1.41
Verify score		-0.18	0.21	0.17	0.3	0.13	0.19	0.08	0.37
PSI- BLAST		3.60E-28	1.80E-27	5.40E-20	1.80E-30	1.80E-50	9.00E-29	1.80E-38	1.80E-32
End		1860	1960	354	2062	2163	2264	2371	2473
Start AA		0691	1780	182	1898	1975	2104	2178	2306
Chain		4	4	¥	¥	¥	4	<	∢
PDB ID		ledh	ledh	ledh	ledh	1edh	ledh	ledh	1edh
SEQ D		808	808	808	808	808	808	808	808

									
PDB annotation	ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELLA CADHERN DOMAINS I AND 2, ECAD 2. ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELLAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECADIS, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELJAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM
Coumpound		E-CADHERIN, CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;
SeqFold score				·	120.62				
PMF		0.86	0.75	0.99		1	1202.08	0.94	0.41
Verify score		0.07	0.14	0.33		0.52	0.46	0.31	0.15
PSI- BLAST		3.60E-29	1.60E-39	7.20E-31	5.40E-58	5.40E-58	1.80E-33	1.80E-28	5.40E-24
End		2577	2683	2789	2895	2898	3003	3105	450
Start AA		2414	2488	2619	2692	2693	2831	2941	296
Chain ID		4	4	4	∢	₹ .	4	∢	A
PDB		1edh	ledh	1edh	ledh	1edh	ledh	1edh	ledh
SEQ NO D		808	808	808	808	808	808	808	808

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PDB annotation	BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD 12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN
Coumpound		E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;
SeqFold score		·							
PMF		9.6	0.59	86.0	-	0.3	0.89	0.77	-
Verify score		0.28	0.15	0.53	0.47	0.12	0.31	0.5	0.33
PSI- BLAST		1.80E-27	1.80E-28	1.40E-35	1.80E-16	3.60E-53	5.40E-30	3.60E-27	1.10E-57
End		3191	3313	3418	3523	248	556	662	917
Start AA		3045	3147	3225	3355	39	401	465	718
Chain		V	4	4	4	∢	4	Ą	V
PDB		1edh	1edh	ledh	Iedh	1edh	ledh	ledh	ledh
SEQ	ž	808	808	808	808	808	808	808	808

PDB annotation	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	MATRIX PROTEIN EXTRACELLULAR MATRIX, CALCIUM-BINDING, GLYCOPROTEIN, 2 REPEAT, SIGNAL, MULTIGENE FAMILY, DISEASE MUTATION, 3 EGF-LIKE DOMAIN, HUMAN FIBRILLIN-1 FRAGMENT, MATRIX PROTEIN	MATRIX PROTEIN EXTRACELLULAR MATRIX, CALCIUM-BINDING, GLYCOPROTEIN, 2 REPEAT, SIGNAL, MULTIGENE FAMILY, DISEASE MUTATION, 3 EGF-LIKE DOMAIN, HUMAN FIBRILLIN-1 FRAGMENT, MATRIX PROTEIN	BLOOD CLOTTING COMPLEX(SERINE PROTEASE/COFACTORLIGAND), BLOOD COAGULATION, 2 SERINE PROTEASE, COMPLEX, CO-FACTOR, RECEPTOR ENZYME, 3 INHIBITOR, GLA, EGF, COMPLEX (SERINE 4 PROTEASE/COFACTOR/LIGAND), BLOOD CLOTTING	BLOOD CLOTTING COMPLEX(SERINE PROTEASE/COFACTOR/LIGAND), BLOOD COAGULATION, 2 SERINE PROTEASE, COMPLEX, CO-FACTOR, RECEPTOR ENZYME, 3 INHIBITOR, GLA, EGF, COMPLEX (SERINE 4 PROTEASE/COFACTOR/LIGAND), BLOOD CLOTTING	GLYCOPROTEIN GLYCOPROTEIN
Coumpound	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	FIBRILLIN; CHAIN: NULL;	FIBRILLIN; CHAIN: NULL;	BLOOD COAGULATION FACTOR VIIA; CHAIN: L; BLOOD COAGULATION FACTOR VIIA; CHAIN: H; SOLUBLE TISSUE FACTOR; CHAIN: T; \$L15; CHAIN: I;	BLOOD COAGULATION FACTOR VILA; CHAIN: L; BLOOD COAGULATION FACTOR VIIA; CHAIN: H; SOLUBLE TISSUE FACTOR; CHAIN: T; 5L15; CHAIN: I;	LAMININ; CHAIN: NULL;
SeqFold score		٠					
PMF	_	0.52	0.48	-0.18	10.0-	-0.19	-0.2
Verify	0.14	0.24	90.0	0.02	0.12	0.2	0.07
PSI- BLAST	1.80E-33	1.80E-30	3.60E-16	9.00E-14	3.60E-14	1.30E-13	1.80E-17
End	1022	1129	3931	4217	3976	4259	4228
Start	854	959	3860	4126	3900	4170	4074
Chain	A	V			ـــ	1	
PDB ID	ledh	ledh	n n	lem n	lfak	1fak	1klo
SEQ D	808	808	808	808	808	808	808

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	TEIN	ACHEK	ADHER DHER	ADHER	ADHER	ADHER	ADHER	ADHER	CADHER	SADHER	ADHER	ADHER	ADHER								
tation	GLYCOPROTEIN GLYCOPROTEIN	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCG (3	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13
PDB annotation	EIN GL	ON PK	ION PRO	ION PRO	ION PR(ION PR ON PR	ION PR	ION PR	ION PR	ION PR	ION PR	ION PR	ION PR	ION PR	ION PR	ION PR					
a	OPROT	ADHES 13	ADHES	ADHES	ADHES	ADHES 13	ADHES 13	ADHES 13	ADHES 13	ADHES 13	ADHES 113	ADHES 113	ADHES 113	ADHES 113	ADHES 13						
	GLYC	CELL AL	CELL AD INCG 13	CELL AD INCG 13	CELL AD	CELL AD	CELL AD INCG 13	CELL AD	CELL AD	CELL AD	CELL AD	CELL AD	CELL AD INCG 13	CELL AL INCG 13	CELL AD INCG 13	CELL AD INCG 13	CELL AD	CELL AD	CELL AD	CELL AE	CELL AD INCG 13
	į.																				
Coumpound	LAMININ; CHAIN: NULL;	N-CADHERIN; 1NCG 3	N-CADHERIN; 1NCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3	N-CADHERIN; 1NCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3
Coum.	IIN; CH	HERIN;	HERIN;	HERIN;	HERIN,	HERIN;	HERIN;	HERIN;	HERIN;	HERIN;	HERIN;	HERIN;	HERIN	HERIN							
	LAMIN	N-CAD -CAD	N-CAD	N-CAD	N-CAD	N-CAD	N-CAD	N-CAD	N-CAL	N-CAL	N-CAL	N-CAL									
SeqFold score																		: :			
PMF score	-0.2	0.63	0.87	0.3	0.72	60.0	0.04	0.31	0.1	0.29	10.0	0.64	0.22	0.25	0.7	0.51	0.34	0.45	8.0	0.53	0.46
Verify	0.07	0.12	0.15	0.34	0	0.15	-0.45	0.35	-0.06	0.31	0.11	0.43	-0.23	0.41	0.44	0.45	90.0	0.28	0.16	-0.19	0.55
PSI- BLAST	1.80E-18	1.60E-05	5.40E-05	3.60E-17	5.40E-05	3.60E-05	9.00E-06	3.60E-06	1.80E-15	0.00036	9.00E-07	1.80E-06	45	0.00014	3.60E-20	9.00E-07	0.00018	1.10E-12	1.10E-21	0.00036	1.60E-05
	╀	-		₩	 		 	├		-	├—		8 0.0045	\vdash	├-	├	├	! —	-	 	1
End	4262	1127	1232	1439	1546	229	1651	1748	2061	2161	2262	2369	2458	2681	2788	3106	3191	3312	811	8	1021
Start AA	4134	1064	1169	1350	1455	155	1599	1667	1975	2079	2180	2304	2411	2592	2692	3039	3120	3225	716	852	932
Chain																					
PDB	1klo	Incg	lncg	lncg	lncg	Incg	Incg	Incg	Incg	Incg	lncg	Incg	lncg	lncg	Incg	lncg	lncg	lncg	Incg	Incg	lncg
SEQ D	808	808	808	808	808	808	808	808	808	808	808	808	808	808	808	808	808	808	808	808	808

PDB annotation	CELL ADHESION PROTEIN CADHERIN INCI 13	CELL ADHESION PROTEIN CADHERIN INCI 13	CELL ADHESION PROTEIN CADHERIN INCI 13	CELL ADHESION PROTEIN CADHERIN INCI 13	CELL ADHESION PROTEIN CADHERIN INCI 13	CELL ADHESION PROTEIN CADHERIN INCI 13	CELL ADHESION PROTEIN CADHERIN INCI 13	CELL ADHESION PROTEIN CADHERIN INCI 13	CELL ADHESION PROTEIN CADHERIN INCI 13	CELL ADHESION PROTEIN CADHERIN INCI 13	CELL ADHESION PROTEIN CADHERIN INCI 13	CELL ADHESION PROTEIN CADHERIN INCI 13	CELL ADHESION PROTEIN CADHERIN INCI 13	CELL ADHESION PROTEIN CADHERIN INCI 13	CELL ADHESION PROTEIN CADHERIN INCI 13	CELL ADHESION PROTEIN CADHERIN INCI 13	CELL ADHESION PROTEIN CADHERIN INCI 13	CELL ADHESION PROTEIN CADHERIN INCI 13	CELL ADHESION PROTEIN CADHERIN INCI 13	CELL ADHESION PROTEIN CADHERIN INC! 13	CELL ADHESION PROTEIN CELL
Coumpound	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3	N-CADHERIN, INCI 3	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3	N-CADHERIN; CHAIN: A;
SeqFold score																					
PMF score	0.78	0.9	90:0	0.16	0.64	0.4	0.27	0.07	0.3	9.0	0.49	-	0	0.78	99.0	0.17	0.17	0.88	0.84	0.59	1
Verify score	90.0	-0.16	0.53	0.21	0.26	-0.17	-0.5	0.24	-0.4	0.39	-0.03	0.56	-0.01	-0.24	0.52	-0.2	0.31	0.01	0.26	0.4	0.39
PSI- BLAST	3.60E-06	1.80E-05	3.60E-16	5.40E-05	1.40E-06	5.40E-05	0.0045	1.80E-14	3.60E-06	5.40E-07	0.0079	1.80E-19	1.80E-06	0.0045	1.10E-06	0.0013	1.40E-10	5.40E-20	7.20E-05	7.20E-06	1.10E-35
End AA	1129	1234	1440	1547	1750	248	1960	2062	2264	2371	2458	2789	2898	3003	3105	3191	3313	812	917	1022	1234
Start AA	1065	1178	1350	1481	1991	181	1912	1975	2180	2307	2412	2692	2832	2951	3041	3146	3225	716	853	932	1039
Chain ID	æ	æ	æ	В	В	æ	Ф	В	В	В	В	В	æ	В	В	В	В	В	В	В	A
PDB ID	Inci	Inci	Inci	- Juci	Inci	Inci	Inci	Inci	Inci	Inci	Inci	Inci	Inci	Inci	Inci	Inci	Inci	Inci	Inci	Inci	Incj
SE SE	808	808	808	808	808	808	808 808	808	808	808	808	808	808	808	808	808	808	808	808	808	808

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PDB annotation	ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
Coumpound		N-CADHERIN: CHAIN: A;	N-CADHERIN: CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;
SeqFold score																			120.25		
PMF score		-	91.0	0.99	0.88	_	0.84	0.23	1	6.0	0.45	68.0	1	1	8.0	9.0	0.51	1		_	0.99
Verify score		0.53	1.0	0.28	0.02	0.46	0.2	0.08	0.31	0.25	0.02	0.23	0.25	0.32	-0.03	90.0	0.28	0.23		0.28	0.28
PSI- BLAST		1.40E-31	1.80E-23	1.60E-58	3.60E-35	1.10E-33	7.20E-33	3.60E-22	9.00E-27	3.60E-31	1.60E-56	9.00E-31	3.60E-40	3.60E-35	9.00E-31	3.60E-40	1.30E-26	1.40E-31	3.60E-63	3.60E-63	9.00E-36
End AA		1338	1440	1547	1652	1750	1861	354	0961	2062	2163	2264	2371	2473	2577	2683	450	2789	2897	2898	3003
Start AA		1144	1251	1351	1455	1562	1991	173	1793	1898	1975	2079	2180	2300	2407	2488	256	2592	2691	2693	2825
Chain ID		A	A	٧	Ą	4	A	A	¥	A	V	۷	٧	4	∢	٧	∢	< −	∢	٧	A
PDB ID		Incj	Incj	lncj	Incj	lncj	Incj	Incj	lncj	Incj	lncj	Incj									
SEQ ID NO:		808	808	808	808	808	808	808	808	808	808	808	808	808	808	808	808	808	808	808	808

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PDB annotation	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	SERINE PROTEASE FVIIĄ; FVIIĄ; BLOOD COAGULATION, SERINE PROTEASE		SERINE PROTEASE FVIIA; FVIIA; BLOOD COAGULATION, SERINE PROTEASE		SERINE PROTEASE FVIIA; BLOOD COAGULATION, SERINE PROTEASE
Coumpound	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	COAGULATION FACTOR VIA (LIGHT CHAIN); CHAIN: L; COAGULATION FACTOR VIA (HEAVY CHAIN);	CHAIN: H; TRIPEPTIDYL INHIBITOR; CHAIN: C;	COAGULATION FACTOR VIIA (LIGHT CHAIN); CHAIN: L; COAGULATION FACTOR	VIIA (HEAVY CHAIN); CHAIN: H; TRIPEPTIDYL INHIBITOR; CHAIN: C;	COAGULATION FACTOR VIIA (LIGHT CHAIN); CHAIN: L; COAGULATION FACTOR
SeqFold score																		
PMF score	_	9.0	0.82	-	-	-0.03	0.21	0.36	96.0	10.0	_	0.88	_	0.57		-0.08		-0.15
Verify score	0.35	0.16	0.35	99.0	9.0	0.35	-0.28	0.08	0.36	-0.13	0.39	0.35	0.48	0.68		0.01	•	0.08
PSI- BLAST	9.00E-30	1.30E-30	1.30E-30	1.80E-39	1.80E-17	5.40E-10	1.10E-59	1.80E-35	7.20E-28	1.80E-25	1.10E-63	1.80E-34	3.60E-34	1.20E-21		3.60E-13		7.20E-11
End	3105	3196	3313	3418	3523	3621	248	929	662	812	917	1022	1129	3947		3976		4168
Start	2915	3039	3141	3225	3349	3473	39	390	467	573	717	827	932	3865		3904		4082
Chain	V	A	∀	Ą	V	V	V	A	∢	∢	4	4	4	7		1		1
PDB	Incj	Incj	1ncj	Incj	Incj	Incj	Incj	lncj	lncj	Incj	lncj	lncj	lncj	1qfk		1qfk		1qfk
SEQ NO D	808	· 808	808	808	808	808	808	808	808	808	808	808	808	808		808		808

PDB annotation		CELL ADHESION UVOMORULIN; CADHERIN, CALCTUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN;
Coumpound	VIIA (HEAVY CHAIN); CHAIN: H; TRIPEPTIDYL INHIBITOR; CHAIN: C;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN;
SeqFold score		•												
PMF score		0.13	0.27	0.45	0.4	0.62	0.25	0.63	0.11	0.07	0.31	0.09	99.0	99.0
Verify score		0.33	0.24	0.04	-0.03	0.54	0.3	0.45	0.14	0.35	0.41	0.2	0.07	90.0
PSI- BLAST		7.20E-08	7.80E-20	1.30E-17	7.20E-08	1.30E-10	1.80E-21	1.30E-08	5.40E-07	2.60E-07	1.00E-12	1.40E-08	1.30E-12	3.60E-06
End		1133	1133	1238	1238	1334	1444	1549	1551	526	1650	1656	1754	1754
Start AA		1026	1041	1145	1711	1249	1350	1455	1455	155	1570	1589	1667	0691
Chain														
PDB D		1suh	1suh	1suh	1suh	1 suh	1suh	1suh	1suh	lsuh	1suh	1suh	1suh	1suh
SEQ Seq		808	808	808	808	808	808	808	808	808	808	808	808	808

S B S	PDB ID	Chain ID	Start AA	End AA	PSI- BLAST	Verify score	PMF	SeqFold score	Conmpound	PDB annotation
										ADHESION
808	1suh		1777	1867	1.30E-14	-0.06	0.48		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	1suh		1780	1868	3.60E-05	-0.24	0.39		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	1suh		182	252	5.40E-07	-0.41	0.13		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	1suh		1898	1964	1.60E-05	-0.1	0.18		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UYOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	1 suh		1975	2066	7.20E-18	0.17	1202.08		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	1suh		2084	2163	2.60E-15	0.42	60.0		EPITHELIÁL CADHERIN; CHAIN: NULL;	CELL ADHESION UYOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	lsuh		2104	2167	3.60E-07	0.05	0.12		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	1suh		2178	2268	1.30E-14	-0.23	0.57		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	1suh		2178	2268	3.60E-10	-0.36	0.46		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UYOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	1 suh		2280	2375	3.90E-20	0.44	0.77		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	1suh		2306	2375	1.10E-08	0.4	0.94		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	Isuh		2395	2475	3.90E-05	0.03	0.58		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN: CADHERIN, CALCIUM BINDING, CELL ADHESION
808	lsuh		2414	2477	0.00011	-0.37	0.06		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	1suh		2488	2581	3.60E-12	-0.2	0.03		EPITHELIAL CADHERIN;	CELL ADHESION UVOMORULIN;

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PDB annotation	CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULN; CADHERN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UYOMORULIN; CADHERN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
Coumpound	CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;
SeqFold									-					
PMF		86:0	0.12	66.0	0.87	0.53	0.93	0.86	0.89	0.55	0.25	0.82	96.0	0.94
Verify score		0.16	0.02	0.51	0.43	-0.01	0.37	-0.07	0.36	0.33	0.43	9.0	0.58	9.0
PSI- BLAST		1.30E-13	2.60E-10	1.40E-23	2.60E-21	1.40E-08	1.20E-14	1.80E-05	2.60E-21	3.60E-06	3.90E-12	.1.30E-13	3.90E-21	5.40E-05
End		2581	2681	2793	2902	2902	3007	3007	3109	3109	3213	3317	3422	3422
Start		2489	2591	2692	2806	2831	2914	2941	3026	3045	3120	3225	3330	3355
Chain ID														·
PDB		lsuh	Isuh	1soh	Isuh	lsuh	1suh	1suh	1suh	1suh	1suh	lsuh	1suh	1suh
SEQ NO:		808	808	808	808	808	808	808	808	808	808	808	808	808

		ERIN; CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION		ERIN; CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	ERIN; CELL ADHESION UVOMORULIN; CADHERIN. CALCIUM BINDING, CELL ADHESION				ERIN; CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION			N. L, C, STUART FACTOR, BLOOD COAGULATION FACTOR COAGULATION FACTOR, SERINE PROTEINASE, EPIDERMAL 2 GROWTH FACTOR I IKE DOMAIN	
Coumpound	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN; NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	T-PLASMINOGEN ACTIVATOR F1-G; 1TPG 7 CHAIN: NULL: 1TPG 8	BLOOD COAGULATION FACTOR XA; CHAIN: L, C;	LECTIN (AGGLUTININ)
SeqFold score								•					_
PMF	9.65	0.24	0.3	0.19	0.57	0.95	0.72	0.47	0.57	0.18	0.59	0.1	-0.2
Verify score	0.56	0.08	-0.21	-0.03	0.39	-0.21	0.4	0.27	0.37	0.2	0.21	-0.26	0.02
PSI- BLAST	6.50E-11	0.0001	9.00E-06	1.30E-15	3.90E-05	7.20E-25	1.00E-17	1.30E-07	5.20E-13	9.00E-07	1.30E-18	7.20E-11	5.40E-12
End	3524	454	454	260	859	816	921	921	1026	1026	3940	3994	4234
Start AA	3435	363	401	467	290	716	826	854	930	656	3863	3904	4069
Chain								,				J	٧
EQE CI	1suh	lsuh	1suh	1suh	1suh	1suh	1suh	1suh	Isuh	Isuh	ltpg	lxka	9wga
SEQ NO B	808	808	808	808	808	808	808	808	808	808	808	808	808

PDB annotation	GROWTH FACTOR (ABU6, 20) MEGF4- 48; GROWTH FACTOR, MURINE EPIDERMAL GROWTH FACTOR, DISULFIDE 2 CONNECTIVITIES, EGF- LIKE DOMAIN, REPEAT	COMPLEX (BLOOD COAGULATION/INHIBITOR) AUTOPROTHROMBIN 11A; HYDROLASE, SERINE PROTEINASE), PLASMA CALCIUM BINDING, 2 GLYCOPROTEIN, COMPLEX (BLOOD COAGULATION/INHIBITOR)	BLOOD COAGULATION BLOOD COAGULATION, EGF, HYDROLASE, SERINE PROTEASE	MEMBRANE PROTEIN LECTIN-LIKE, NEUROBIOLOGY, CELL-CELL ADHESION, CELL-CELL 2 RECOGNITION, ALTERNATIVE SPLICING, MEMBRANE PROTEIN	TRANSPORT PROTEIN SHBG; STEROID TRANSPORT, LAMININ G- LIKE DOMAIN. JELLYROLL, 2 ANDROGEN BINDING PROTEIN (ABP), SEX STEROID BINDING PROTEIN 3 (SBP)	HYDROLASEHYDROLASE INHIBITOR PROTEIN-PEPTIDE COMPLEX	HYDROLASEHYDROLASE INHIBITOR PROTEIN-PEPTIDE COMPLEX
Coumpound	EPIDERMAL GROWTH FACTOR; CHAIN: NULL;	ACTIVATED PROTEIN C; CHAIN: C, L; D-PHE-PRO- MAI; CHAIN: P;	FACTOR VII; CHAIN: NULL;	NEUREXIN-I BETA; CHAIN: A, B, C, D, E, F, G, H;	SEX HORMONE-BINDING GLOBULIN; CHAIN: A;	DES-GLA FACTOR VIIA (HEAVY CHAIN); CHAIN: H, I; DES-GLA FACTOR VIIA (LIGHT CHAIN); CHAIN: L, M; (DPN)-PHE-ARG; CHAIN: C, D; PEPTIDE E-76; CHAIN: X, Y;	DES-GLA FACTOR VIIA (HEAVY CHAIN); CHAIN: H, I; DES-GLA FACTOR VIIA (LIGHT CHAIN); CHAIN: L, M; (DPN)-PHE-ARG; CHAIN: C, D; PEPTIDE E-76; CHAIN: X, Y;
SeqFold score							·
PMF .	0.58	-0.12	0.99	0.4	0.72	0.05	-0.18
Verify	0.39	0.55	96.0	0.26	0.46	0.48	0.25
PSI- BLAST	6.50E-12	6.50E-19	1.30E-11	1.30E-26	2.60E-27	3.90E-21	1.30E-13
End AA	3984	4033	3987	3928	3934	4033	4344
Start AA	3949	3943	3949	3771	3773	3944	4263
Chain ID		1		4	∢	Г	1
PDB ID	la3p	laut	1669	lodr	142s	ldva	ldva
SEQ NO:	808	808	608	809	809	808	809

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PDB annotation	SERINE PROTEINASE COAGULATION FACTOR II; COAGULATION FACTOR II; FETOMODULIN, TM, CD141 ANTIGEN; EGR-CMK SERINE PROTEINASE, EGF-LIKE DOMAINS, ANTIFIBRINOLYTIC COMPLEX ANTIFIBRINOLYTIC COMPLEX	CELL ADHESION PROTEIN EPITHELLAL CADHERIN DOMAINS 1 AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN, CALCIUM	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECADIZ, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECADI 2, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN; CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECADIE; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12; CADHERIN, CELL
Coumpound	THROMBIN LIGHT CHAIN; CHAIN; A, B, C, D; THROMBIN HEAVY CHAIN; CHAIN; M, N, O, P; THROMBOMODULN; CHAIN; I, J, K, L; THROMBIN DMIBITOR L-GLU-L-GLY-L-ARM; CHAIN; E, F, G, H;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;
SeqFold score								
PMF	-0.19 ·	p-4	1202.08	6:0	0.82	-1.41	96.0	0.99
Verify	0.06	0.26	0.22	0.45	0.27	0.1	0.21	0.19
PSI- BLAST	1.30E-11	3.60E-33	1.30E-32	3.605-21	1.605-49	1.30E-32	3.60E-33	1.60E-28
End	4336	1234	1338	1440	1547	1652	1750	1860
Start	4232	1066	1171	1279	1352	1460	1589	1690
Chain	I	V	4	∢	4	⋖	. 4	4
PDB	IdxS	1edh	ledh	1edh	ledh	ledh	1edh	Igh H
SEQ	808 809	809	808	608	808	809	808	809

PDB annotation	ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECADIS; CADHERIN, CELL ADHESION PROTEIN, CALCIUM HNDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM
Coumpound	·	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAÎN: A, B;
SeqFold score									,
PMF		-	0	0.96	0.51	0.78	-	1	0.86
Verify score		0.32	0	0.26	0.3	0.24	0.18	0.32	0.07
PSI- BLAST		3.60E-26	1.60E-20	3.60E-29	1.10E-50	1.30E-28	3.60E-48	1.60E-35	1.80E-29
End AA		0961	354	2062	2163	2264	2371	2473	2577
Start AA		1800	182	1898	1975	2104	2178	2306	2414
Chain		<	∢	∢	∢	∢	¥	∢	∢
PDB ID		1edh	ledh	ledh	ledh	1edh	ledh	1cdh	ledh
SEQ EQ		608	608	608	608	608	608	808	808

PDB annotation	BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM ENEMING DE OFFEN	BINDING FINGERIA	CELL ADRESION PROTEIN EPITHELIAL CADHERIN DOMAINS I	AND 2, ECAD12; CADHERIN, CELL	ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN	EPITHELIAL CADHERIN DOMAINS I	AND 2, ECAD12; CADHERIN, CELL	ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN	EPITHELIAL CADHERIN DOMAINS I	AND 2, ECAD12; CADHERIN, CELL	BINDING PROTEIN	CELL ADHESION PROTEIN	EPITHELIAL CADHERIN DOMAINS I	AND 2, ECAD12; CADHERIN, CELL	ADHESION PROTEIN, CALCIUM	BINDING PROTEIN	CELL ADHESION PROTEIN	EPITHELIAL CADHERIN DOMAINS I	AND 2, ECADI2; CADHERIN, CELL	ADHESION PROTEIN, CALCIUM	BINDING PROTEIN	CELL ADHESION PROTEIN EPITHEL 141 CADHEBIN DOMAINS 1	AND BOADS CADIFERN CELL	AND 2, ECADIZ; CADRENIN, CELL	ADHESION PROTEIN, CALCIUM	CELL ADHESTON PROTEIN	EPITHELIAL CADHERIN DOMAINS 1	AND 2, ECAD12; CADHERIN, CELL	ADHESION PROTEIN, CALCIUM BINDING PROTEIN
Conmpound		E-CADHERIN; CHAIN: A, B;		E-CADHERIN; CHAIN: A, B;			E-CADHERIN: CHAIN: A. B.				E-CADHERIN; CHAIN: A, B;				E-CADHERIN; CHAIN: A, B;					E-CADHERIN; CHAIN: A, B;					E-CADHERIN; CHAIN: A, B;				E-CADHERDS CHAIN: A B:	, c. (1) (1) (1) (1) (1) (1) (1) (1) (1) (1)		
SeqFold score							120.62																									
PMF		0.93		96.0							_				-	,				0.94					0.53				0 64	5		
Verify score		0		0.1							0.47				0.27					0.31					0.38				0.16	2		
PSI- BLAST		1.80E-38		5.40E-32			1 80E-57	200:1			1.80E-57				7.20E-35					5.40E-29					7.20E-25				S 40E 38	97-704-6		
End		2683		2789			2000	7657			2898				3003					3105					450				2107	212/		
Start		2488		5619			2607	7037			2693			_	2831	<u> </u>				2941					536	_	_		2046	2010		
Chain		¥		٧				ť			A				 	:				4					∢				<	<		
PDB	1	1edh		1edh			1	III			Iedh	_			ledh					ledh					ledh				1 adh	100		
SEQ U	ÿ	808		808			8	6			809				808	}				808					60g				000	600		

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PDB annotation	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELLA CADHERIN DOMAINS 1 AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD 12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN
Coumpound	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;
SeqFold						-			
PMF	-1.41	0.98	1	0.3	0.83	0.82	0.27	-	_
Verify	0.57	0.45	0.66	0.12	0.29	0.17	-0.05	0.33	0.31
PSI- BLAST	1.80E-32	1.80E-48	3.60E-30	1.80E-51	1.10E-29	3.60E-29	5.40E-22	1.10E-55	1.10E-32
End	3313	3418	3523	248	556	662	812	917	1022
Start AA	3120	3225	3355	39	406	464	591	718	854
Chain 10	V	∢	<	<	<	4	٧	٧	٧
PDB TD	1edh	ledh	ledh	1edh	ledh	1edh	ledh	ledh	1cdh
SEQ EQ	808	808	608	608	608	808	808	808	809

PDB annotation					GLYCOPROTEIN GLYCOPROTEIN	GLYCOPROTEIN GLYCOPROTEIN	GLYCOPROTEIN GLICOFROTEIN	INCG 13	CELL ADHESION PROTEIN CADHERIN	CELL ADHESION PROTEIN CADHERIN 1NCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13
Coumpound		E-CADHERIN; CHAIN: A, B;	FIBRILLIN; CHAIN: NULL;	P-SELECTIN; CHAIN: NULL;	LAMININ; CHAIN: NULL;	LAMININ; CHAIN: NULL;	LAMININ; CHAIN: NULL;	N-CADHERIN; INCG 3	N-CADHERIN; 1NCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3
SeqFold																
PMF		69.0	0.99	0.88	0.27	-0.03	-0.2	0.29	69.0	0.29	1.0	0.09	0.09	0.31	0.1	0.11
Verify		0.11	0.71	1.03	9.1	0.07	0.23	0.12	0.13	0.01	0.46	0.15	-0.13	0.35	0.22	0.28
PSI- BLAST		9.00E-32	1.80E-16	1.30E-11	1.10E-13	5.40E-21	3.60E-17	9.00E-06	0.00014	1.40E-14	0.00018	3.60E-05	9.00E-06	3.60E-06	3.60E-17	0.00036
End		1129	4022	3988	4050	4075	4342	1127	1232	1439	1546	229	1650	1748	2061	2161
Start AA		959	3946	3949	3924	3954	4201	1062	1167	1350	1480	155	1599	1667	1970	2079
Chain		<			-										_	_
PDB		ledh	lem n	1£b	1660	Iklo	1K9	. Incg	lncg	Incg	Incg	Incg	Incg	Incg	lncg	Incg
SEQ	Ö	808	809	808	800	8	808	808	809	808	809	809	808	808	808	808

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PDB annotation	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN	CELL ADHESION PROTEIN CADHERIN INCI 13	CELL ADHESION PROTEIN CADHERIN INCI 13	CELL ADHESION PROTEIN CADHERIN INCI 13	CELL ADHESION PROTEIN CADHERIN INCI 13	CELL ADHESION PROTEIN CADHERIN INCI 13	CELL ADHESION PROTEIN CADHERIN	CELL ADHESION PROTEIN CADHERIN	CELL ADHESION PROTEIN CADHERIN
Соитроипд	N-CADHERIN; INCG 3	N-CADHERIN; 1NCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3
SeqFold score																					
PMF score	0.1	0.74	0.43	0.57	0.7	0.39	0.51	0.34	0.36	8.0	0.53	0.64	0.78	0.96	0.22	0.65	9.0	0.4	0.23	0.05	0.1
Verify score	-0.08	0.41	-0.27	0.34	0.44	0.35	0.45	90.0	0.33	0.16	-0.19	0.38	90.0	90.0	0.43	-0.08	0.35	-0.17	0.08	-0.25	-0.11
PSI- BLAST	3.60E-12	1.80E-06	0.00036	1.60E-05	1.80E-19	0.00054	3.60E-06	1.80E-05	1.60E-11	7.20E-20	0.00018	5.40E-05	1.80E-06	5.40E-05	1,30E-13	0.00018	9.00E-07	1.60E-05	1.80E-16	0.0013	1.30E-11
End AA	2263	2370	2458	2681	2788	2988	3106	3191	3311	811	006	1003	1129	1234	1440	1547	1750	248	2902	2163	2264
Start AA	2178	2304	2411	2593	2692	2913	3039	3120	3225	716	852	932	1065	1172	1350	1491	1667	181	1970	2116	2178
Chain 10													æ	В	В	B	gg	B.	В	В	В
aga au	lncg	Incg	lncg	Incg	Incg	Incg	lncg	lncg	lncg	Boul	Jucg	Incg	Inci	1nci	Inci	Inci	Inci	lnci .	Inci	Inci	Inci
SEQ NO:	809	608	808	808	808	809	809	808	608	809	608	608	808	608	809	808	808	808	808	809	608

PDB annotation	INCI 13	CELL ADHESION PROTEIN CADHERIN	CELL ADHESION PROTEIN CADHERIN	CELL ADHESION PROTEIN CADHEKIN	CELL ADHESION PROTEIN CADHERIN	CELL ADHESION PROTEIN CAUHERIN INCI 13													-		N: A; CELL ADHESION PROTEIN CELL ADHESION PROTEIN
Coumpound		N-CADHERIN; INCI 3	N-CADHERIN; INCI 3	N-CADHERIN; 1NCI 3	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;
SeqFold score																					
PMF		0.75	0.58	90.0	-	0.92	0.42	0.17	0.49	0.99	96.0	0.71	0.63	_	-	0.54	0.99	-	0.21		0.74
Verify		0.17	-0.14	0	95.0	0.42	0.56	-0.2	19.0	0.87	0.15	-0.4	0.36	0.45	0.34	0.3	0.29	0.03	0.05	0.45	-0.05
PSI- BLAST		5.40E-07	0.0009	1.60E-05	3.60E-19	0.0036	5.40E-06	0.00036	1.80E-10	1.80E-08	1.80E-19	5.40E-05	1.80E-05	3.60E-36	5.40E-33	1.80E-22	3.60E-53	1.80E-34	5.40E-24	3.60E-33	3.60E-32
End		2371	2458	2683	2789	3003	3105	3191	3313	3418	812	917	1022	1234	1338	1440	1547	1652	354	1750	1861
Start		2307	2414	2620	2692	2942	3044	3146	3225	3354	715	862	932	1039	1147	1270	1351	1458	155	1562	1667
Chain		В	В	В	В	В	В	В	æ	m	8	B	В	A	4	4	4	<	<	<	4
PDB	1	Inci	Inci	lnci	Inci	Inci	Inci	Inci	Inci	Inci	Inci	Inci	1nci	Incj	1ncj	Incj	Incj	Incj	Incj	Incj	lncj
SEQ	Ö	809	809	809	809	809	809	808	809	809	808	809	808	808	808	809	809	809	808	808	808

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PDB annotation	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL
Coumpound	N-CADHERIN; CHAIN: A;	N-CADHERIN: CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN: CHAIN: A:
SeqFold score											120.25										
PMF	. 96'0	0.76	0.33	0.86	-	1	0.75	0.83	0.3	-			66.0		0.58	0.63	0.99	-	0.39	0.21	0.35
Verify score	0.31	0.42	0.11	0.48	0.17	0.27	-0.1	0.14	0.25	0.29		0.28	0.31	0.62	-0.03	0.36	0.48	0.63	0.52	-0.28	0.3
PSI- BLAST	7.20E-27	3.60E-28	7.20E-55	3.60E-30	5.40E-52	5.40E-36	7.20E-32	1.60E-41	3.60E-27	1.30E-32	5.40E-63	5.40E-63	3.60E-38	1.80E-29	1.40E-31	1.80E-34	7.20E-51	3.60E-32	3.60E-13	1.80E-57	1.80E-34
End	1960	2062	2163	2264	2371	2473	2577	2683	450	2789	2897	2898	3003	3105	3196	3313	3418	3523	3621	248	556
Start AA	1782	1898	1970	2079	2178	2300	2407	2488	256	2593	2691	2693	2825	2913	3039	3120	3225	3346	3433	39	390
Chain TD	A	A	A	<	∢	<	V V	▼	<	A	⋖	¥	V	<	4	∢	<	⋖	4	4	×
PDB	Incj	Incj	Incj	lncj	lncj	1ncj	1ncj	Incj	Incj	1ncj	Incj	Incj	Incj	Jucj	1ncj	Incj	Incj	Incj	Incj	Incj	Inci
SEQ	Ö 808	809	608	809	608	608	608	608	608	809	809	809	809	809	608	809	608	808	808	608	808

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PDB annotation	ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	COMPLEX (BLOOD COGGULATION/INHIBITOR) CHRISTMAS FACTOR; COMPLEX, INHIBITOR, HEMOPHILIA/EGF,	BLOOD COAGULATION, 2 PLASMA, SERINE PROTEASE, CALCIUMBINDING, HYDROLASE, 3 GLYCOPROTEIN	SERINE PROTEASE FVIIA: BLOOD COAGULATION, SERINE PROTEASE	SERINE PROTEASE FVIIA; FVIIA; BLOOD COAGULATION, SERINE PROTEASE	SERINE PROTEASE FVIIA; FVIIA; BLOOD COAGULATION, SERINE PROTEASE	SERINE PROTEASB FVIIA; BLOOD COAGULATION, SERINE PROTEASE
Coumpound		N-CADHERIN; CHAIN: A;	FACTOR IXA; CHAIN: C, L,; D-PHE-PRO-ARG; CHAIN: I;		COAGULATION FACTOR VIIA (LIGHT CHAIN); CHAIN: L; COAGULATION FACTOR VIIA (HEAVY CHAIN); CHAIN: H; TRIPEPTIDYL INHIBITOR: CHAIN: C.	COAGULATION FACTOR VIIA (LIGHT CHAIN); CHAIN: L; COAGULATION FACTOR VIIA (HEAVY CHAIN); CHAIN: H; TRIPEPTIDYL INHIBITOR: CHAIN: C	COAGULATION FACTOR VIIA (LIGHT CHAIN); CHAIN: L; COAGULATION FACTOR VIIA (HEAVY CHAIN); CHAIN: H; TRIPEPTIDYL NHIBITOR: CHAIN: C	COAGULATION FACTOR VIIA (LIGHT CHAIN); CHAIN: 1; COAGULATION FACTOR VIIA (HEAVY CHAIN);				
SeqFold score												
PMF		66.0	0.18	_	0.88	1	-0.19		-0.19	0.57	-0.17	-0.17
Verify score		0.37	0.02	0.23	0.34	0.39	0.01		0.2	0.68	0	0.37
PSI- BLAST		9.00E-30	1.80E-25	7.20E-62	1.80E-34	1.10E-34	1.40E-10		1.30E-11	1.20E-21	5.40E-14	7.20E-13
End		299	812	917	1022	1129	4312		4001	4033	4202	4344
Start		467	112	717	827	932	4224		3943	3951	4121	4267
Chain ID		V	¥	<	¥	A	<u>ـــ</u>		13	ы	L)	7
PDB		Incj	Incj	1ncj	Incj	Incj	lpfx		1qfk	1qfk	1qfk	1qfk
SEQ	ÿ	809	608	608	608	608	608		808	808	608	808

PDB annotation		COAGULATION FACTOR SERINE PROTEINASE, BLOOD COAGULATION, COAGULATION FACTOR	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UYOMOKULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UYOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UYOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UYOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CADHERIN, CALCIUM BINDING, CELL			
Coumpound	CHAIN: H; TRIPEPTIDYL INHIBITOR; CHAIN: C;	COAGULATION FACTOR IX; CHAIN: A; COAGULATION FACTOR IX; CHAIN: B;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;
SeqFold														
PMF score		-0.19	0.27	0.93	0.45	0.62	0.13	0.04	0.63	0.07	0.31	0.13	99.0	0.75
Verify score		0.03	0.24	-0.06	, 40.0	0.54	0.28	0.37	0.45	0.35	0.41	-0.2	0.07	0.17
PSI- BLAST	-	7.80E-14	7.80E-20	1.60E-07	1.30E-17	1.30E-10	0.0013	1.80E-19	1.30E-08	2.60E-07	1.00E-12	9.00E-09	1.30E-12	3.60E-06
End		4001	1133	1133	1238	1334	1342	1444	1549	226	1650	1656	1754	1754
Start AA		3951	1041	1066	1145	1249	1279	1350	1455	155	1570	1589	1667	1690
Chain		В												
PDB		î.	Isuh	1suh	1suh	1suh	Isuh	1suh	Isuh	1suh	1suh	1 suh	1suh	1sth
SEQ	Ö	608	809	809	608	809	809	608	608	608	808	808	809	608

										PDB annotation
[교 ~	PDB C	Chain	Start	AA End	PSI- BLAST	Verify score	PMF	SeqFold	Coumpound	
' I	-+									ADHESION
12	lsuh		1777	1867	1.30E-14	-0.06	0.48		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UYOMOKULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
1	1suh		1800	1868	3.60E-05	-0.18	0.45		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCTUM BINDING, CELL ADHESION
==	1suh		182	252	1.80E-06	-0.41	0.13		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
	1suh		2084	2163	2.60E-15	0.42	0.09		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION O'OMOROLIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
I	1suh		2178	2268	1.30E-15	-0.37	0.4		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN: CADHERIN, CALCIUM BINDING, CELL ADHESION
1-	1suh		2280	2375	3.90E-20	0.44	0.77		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
1	lsuh		2306	2375	3.60E-09	0.35	0.94		· EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UYUMOKULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
1	lsuh		2395	2475	3.90E-05	0.03	0.58		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
1	1suh		2414	2477	3.60E-05	-0.04	0.25		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION DYOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
1.	1suh		2488	2581	1.80E-12	-0.37	0.15		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UYOMOKULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
1	1suh		2489	2581	1.30E-13	0.16	0.98		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UYOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
1	Isuh		2591	2681	2.60E-10	0.02	0.12		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMUKULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
	1suh		2692.	2793	3.60E-23	0.42	0.99		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
+-	Isuh		2806	2902	2.60E-21	0.43	0.87		EPITHELIAL CADHERIN;	CELL ADHESION UVOMORULIN;

PDB annotation	CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UYOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UYOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UYOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UYOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UYOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UYOMORULN; CADHERIN, CALCIUM BINDING, CELL ADHESION
Coumpound	CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;
SeqFold score														
PMF score		0.25	0.93	98.0	0.89	0.72	0.28	0.25	0.82	0.98	0.93	0.65	0.24	0.19
Verify score		0.01	0.37	-0.07	0.36	0.4	0.1	0.43	0.6	0.58	0.66	0.56	0.08	-0.6
PSI- BLAST		3.60E-09	1.20E-14	1.80E-06	2.60E-21	3.60E-06	0.0013	3.90E-12	1.10E-14	3.90E-21	1.30E-09	6.50E-11	0.0001	1.60E-05
End		2902	3007	3007	3109	3109	3185	3213	3317	3422	3422	3524	454	454
Start		2831	2914	2941	3026	3046	3120	3120	3225	3330	3355	3435	363	406
Chain 10										,				
PDB ID		1suh	1suh	1suh	. Isuh	1 suh	1suh	Isuh	lsuh	1suh	1suh	1 suh	1suh	1 suh
SEQ EQ		809	809	608	808	808	809	809	809	808	809	808	608	608

											
PDB annotation	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UYOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	PLASMINOGEN ACTIVATION	BLOOD COAGULATION FACTOR STUART FACTOR; BLOOD COAGULATION FACTOR, SERINE PROTEINASE, EPIDERMAL 2 GROWTH FACTOR LIKE DOMAIN		COMPLEX (BLOOD COAGULATION/MHBITOR) AUTOPROTHROMBIN IIA; HYDROLASE, SERINE PROTEINASE), PLASMA CALCIUM BINDING, 2 GLYCOPROTEIN, COMPLEX (BLOOD COAGULATION/MHBITOR)	BLOOD COAGULATION, SERINE PROTEASE, COMPLEX, CO-FACTOR, 2 RECEPTOR ENZYME, INHIBITOR, GLA, EGF, 3 COMPLEX (SERINE PROTEASE/COFACTOR/LIGAND)
Coumpound	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	T-PLASMINOGEN ACTIVATOR F1-G; 1TPG 7 CHAIN: NULL; 1TPG 8	BLOOD COAGULATION FACTOR XA; CHAIN: L, C;	LECTIN (AGGLUTININ) WHEAT GERM AGGLUTININ (ISOLECTIN 2) 9WGA 3	ACTIVATED PROTEIN C; CHAIN: C, L; D-PHE-PRO- MAI; CHAIN: P;	BLOOD COAGULATION FACTOR VIIA; CHAIN: L, H; SOLUBLE TISSUE FACTOR; CHAIN: T, U; D-PHE-PHE- ARG-
SeqFold score											
PMF	0.19	0.57	0.95	0.72	0.45	0.57	0.59	-0.19	-0.2	-0.12	0.07
Verify score	-0.03	0.39	-0.21	0.4	0.17	0.37	0.21	0.15	0.05	0.55	0.09
PSI- BLAST	1.30E-15	3.90E-05	1.40E-23	1.00E-17	5.40E-07	5.20E-13	1.30E-18	5.40E-12	3.60E-10	6.50E-19	1.40E-12
End	995	658	816	921	921	1026	4026	4348	4337	3947	3897
Start	467	290	716	826	854	930	3949	4267	4166	3857	3831
Chain								l)	∢	 -	J
PDB	1suh	lsuh	lsuh	Isuh	1suh	1suh	1tpg	lxka	9wga	laut	1dan
SEQ EQ	608	608	809	809	809	809	608		809	608	608

PDB annotation		BLOOD COAGULATION, SERINE PROTEASE, COMPLEX, CO-FACTOR, 2 RECEPTOR ENZYME, INHIBITOR, GLA, EGF, 3 COMPLEX (SERINE PROTEASE/COFACTOR/LIGAND)	HYDROLASE/HYDROLASE INHIBITOR PROTEIN-PEPTIDE COMPLEX	HYDROLASE/HYDROLASE INHIBITOR PROTEIN-PEPTIDE COMPLEX	HYDROLASE/HYDROLASE INHIBITOR PROTEIN-PEPTIDE COMPLEX	HYDROLASE/HYDROLASE INHIBITOR PROTEIN-PEPTIDE COMPLEX	HYDROLASE/HYDROLASE INHIBITOR PROTEIN-PEPTIDE COMPLEX
Coumpound	CHLOROMETHYLKETONE (DFFRCMK) WITH CHAIN: C;	BLOOD COAGULATION FACTOR, VIIA; CHAIN: I, H; SOLUBLE TISSUE FACTOR; CHAIN: T, U; D-PHE-PHE- ARG- CHLOROMETHYLKETONE CHERCAKY, WITH CHAIN: C.	DES-GLA FACTOR VIIA (HEAVY CHAIN); CHAIN: H, I; DES-GLA FACTOR VIIA (LIGHT CHAIN); CHAIN: L, M; (DPN)-PHE-ARG; CHAIN: C, D; PEPTIDE E-76; CHAIN: X, Y:	DES-GLA FACTOR VIIA (HEAVY CHAIN); CHAIN: H, I; DES-GLA FACTOR VIIA (LIGHT CHAIN); CHAIN: L, M; (DPN)-PHE-ARG; CHAIN: C, D; PEPTIDE E-76; CHAIN: X, Y;	DES-GLA FACTOR VIIA (HEAVY CHAIN); CHAIN: H, I; DES-GLA FACTOR VIIA (LIGHT CHAIN); CHAIN: L, M; (DPN)-PHE-ARG; CHAIN: C, D; PEPTIDE E-76; CHAIN: X, Y;	DES-GLA FACTOR VIIA (HEAVY CHAIN); CHAIN: H, I; DES-GLA FACTOR VIIA (LIGHT CHAIN); CHAIN: L, M; (DPN)-PHE-ARG; CHAIN: C, D; PEPTIDE E-76; CHAIN: X, Y;	DES-GLA FACTOR VIIA (HEAVY CHAIN); CHAIN: H. I; DES-GLA FACTOR VIIA (LIGHT CHAIN); CHAIN: L.
SeqFold score							
PMF		0.07	0.15	0.05	-0.07	-0.15	-0.19
Verify score		0.1	-0.08	0.48	0.12	0.05	0.25
PSI- BLAST		3.60E-14	1.40E-12	3.90E-21	3.60E-14	3.60E-12	1.30E-13
End		3976	3897	3947	3976	4168	4259
Start		3900	3831	3858	3900	4078	4170
Chain 19		1	ı.	ר	ı	ī	ــــــــــــــــــــــــــــــــــــــ
PDB ID		ldan	Idva	Idva	Idva	Idva	1dva
SEQ P		808	608	608	608	608	608

			- "					·	ī
PDB annotation		CELL ADHESION PROTEIN EPITHELLAL CADHERIN DOMAINS I AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN BPITHELLAL CADHERIN DOMAINS 1 AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECADIS: CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12, CADHERIN, CELL
Coumpound	M; (DPN)-PHE-ARG; CHAIN: C, D; PEPTIDE E-76; CHAIN: X, Y;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAÎN: A, B;	E-CADHERIN; CHAIN: A, B;
SeqFold score								·	
PMF score		.	0.96	0.93	0.89	0.65	-1.41	0.25	0.99
Verify score		0.35	0.16	0.23	0.3	0.03	0.19	-0.18	0.21
PSI- BLAST		1.10E-33	9.00E-30	1.60E-20	3.60E-54	7.20E-32	3.60E-33	3.60E-28	1.80E-27
End AA		1234	1338	1440	1547	1652	1750	1860	1960
Start		1026	1171	1279	1352	1455	1589	1690	1780
Chain		4	4	∢	∢	∢	∀	∢	A
PDB D		1edh	ledh	1edh	ledh	ledh	1edh	ledh	ledh
SEQ SO B		608	808	808	608	809	. 608	809	608

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PDB annotation	ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECADHS, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECADIS; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM
Coumpound		E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;						
SeqFold score									
PMF		0.07	0.94	0.34	0.92	-	-1.41	0.86	0.75
Verify score		0.17	0.3	0.13	0.19	0.08	0.37	0.07	0.14
PSI- BLAST		5.40E-20	1.80E-30	1.80E-50	9.00E-29	1.80E-38	1.80E-32	3.60E-29	1.60E-39
End		354	2062	2163	2264	2371	2473	2577	2683
Start AA		182	1898	1975	2104	2178	2306	2414	2488
Chain		¥.	₹	¥	¥.	∢	∢	· 4	A.
PDB CD		ledh	ledh						
SEQ DO		608	608	808	808	608	808	608	608

PDB annotation	BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECADIS, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12: CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELLAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN
Coumpound		E-CADHERIN; CHAIN: A, B;							
SeqFold score			120.62						
PMF		0.99		_	1202.08	0.94	0.41	9.0	0.59
Verify score		0.33		0.52	0.46	0.31	0.15	0.28	0.15
PSI- BLAST		7.20E-31	5.40E-58	5.40E-58	1.80E-33	1.80E-28	5.40E-24	1.80E-27	1.80E-28
End		2789	2895	2898	3003	3105	450	3191	3313
Start AA		2619	2692	2693	2831	2941	296	3045	3147
Chain		4	<	4	A	A	4	∢	4
PDB CD	1	1edh	ledh	ledh	ledh .	1edh	1edh	ledh	1edh
SEQ EQ	ž	808	608	608	608	608	808	808	809

PDB annotation	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	MATRIX PROTEIN EXTRACELLULAR
Coumpound	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	FIBRILLIN; CHAIN: NULL;
SeqFold score									
PMF	86.0	-	0.3	0.89	0.77	-		0.52	0.48
Verify score	0.53	0.47	0.12	0.31	0.5	0.33	0.14	0.24	90.0
PSI- BLAST	1.40E-35	1.80E-16	3.60E-53	5.40E-30	3.60E-27	1.10E-57	1.80E-33	1.80E-30	3.60E-16
End AA	3418	3523	248	556	662	917	1022	1129	3931
Start	3225	3355	39	401	465	718	854	959	3860
Chain 10	∢	. <	∢	∢	V	<	∢	<	
PDB UD	1edh	ledh .	1edh	ledh .	1edh	1edh	ledh	1cdh	lem
SEQ EQ	608	608	608	809	809	808	808	809	608

PDB annotation	MATRIX, CALCIUM-BINDING, GLYCOPROTEIN, 2 REPEAT, SIGNAL, MULTIGENE FAMILY, DISEASE MUTATION, 3 EGF-LIKE DOMAIN, HUMAN FIBRILLIN-1 FRAGMENT, MATRIX PROTEIN	MATRIX PROTEIN EXTRACELLULAR MATRIX, CALCIUM-BINDING, GLYCOPROTEIN, 2 REPEAT, SIGNAL, MULTIGENE FAMILY, DISEASE MUTATION, 3 EGF-LIKE DOMAIN, HUMAN FIBRILLIN-I FRAGMENT, MATRIX PROTEIN	BLOOD CLOTTING COMPLEX(SERINE PROTEASE/COFACTOR/LIGAND), BLOOD COAGULATION, 2 SERINE PROTEASE, CO-FACTOR, RECEPTOR ENZYME, 3 INHIBITOR, GLA, EGF, COMPLEX (SERINE 4 PROTEASE/COFACTOR/LIGAND), BLOOD CLOTTING	BLOOD CLOTTING COMPLEX(SERINE PROTEASE/COFACTOR/LIGAND), BLOOD COAGULATION, 2 SERINE PROTEASE, COMPLEX, CO-FACTOR, RECEPTOR ENZYME, 3 INHIBITOR, GLA, EGF, COMPLEX (SERINE 4 PROTEASE/COFACTOR/LIGAND), BLOOD CLOTTING	GLYCOPROTEIN GLYCOPROTEIN	GLYCOPROTEIN GLYCOPROTEIN	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13
Coumpound		FIBRILLIN; CHAIN: NULL;	BLOOD COAGULATION FACTOR VIIA; CHAIN: L; BLOOD COAGULATION FACTOR VIIA; CHAIN: H; SOLUBLE TISSUE FACTOR; CHAIN: T; 5L15; CHAIN: I;	BLOOD COAGULATION FACTOR VIIA; CHAIN: L; BLOOD COAGULATION FACTOR VIIA; CHAIN: H; SOLUBLE TISSUE FACTOR; CHAIN: T; 5L15; CHAIN: I;	LAMININ; CHAIN: NULL;	LAMININ; CHAIN: NULL;	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3
SeqFold score											
PMF score		60.18	-0.01	-0.19	-0.2	-0.2	0.63	0.87	0.3	0.72	0.09
Verify score		0.02	0.12	0.2	0.07	0.07	0.12	0.15	0.34	0	0.15
PSI- BLAST		9.00E-14	3.60E-14	1,30E-13	1.80E-17	1.80E-18	1.60E-05	5.40E-05	3.60E-17	5.40E-05	3.60E-05
End		4217	3976	4259	4228	4262	1127	1232	1439	1546	229
Start		4126	3900	4170	4074	4134	1064	1169	1350	1455	155
Chain D			ı	ı,							
PDB ID	c	n n .	·Ifak	Ifak	· 1klo	1klo	Incg	lncg	Incg	lncg	Incg
SEQ B		608	608	608	809	809	809	808	809	808	608

-	Start End	End	-	9	PSI-	Verify	PMF	SeqFold	Coumpound	PDB annotation
ID AA AA BLASI SCOL	AA AA DLASI SUUC	AA DLASI Stute	DLASI SUIC	arone	\Box		200		N.CADHERIN: INCG 3	CELL ADHESION PROTEIN CADHERIN
-0.43	1651 9.00E-06 -0.43	1651 9.00E-06 -0.43	9.00E-06 -0.43	-0.43		ا خ	.		N CADIFFRM: INCC 1	INCG 13 CELL ADHESION PROTEIN CADHERIN
Incg 1667 1748 3.60E-06 0.35 0.31	1748 3.60E-06 0.35	1748 3.60E-06 0.35	3.60E-06 0.35	0.35		0.3			N-CADHEKIN; INCG 3	INCELE ADJUSTICE OF THE BIN OF TH
lncg 1975 2061 1.80E-15 -0.06 0.1	2061 1.80E-15 -0.06	2061 1.80E-15 -0.06	1.80E-15 -0.06	-0.06	-	ö			N-CADHERIN; INCG 3	CELL ADRESION FROTEIN CADRESIN
lncg 2079 2161 0.00036 0.31 0.	2161 0.00036 0.31	2161 0.00036 0.31	0.00036 0.31	0.31	 	o	0.29		N-CADHERIN; 1NCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
lncg 2180 2262 9.00E-07 0.11	2262 9.00E-07 0.11	2262 9.00E-07 0.11	9.00E-07 0.11	0.11		1	0.01		N-CADHERIN; 1NCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
Incg 2304 2369 1.80E-06 0.43	2369 1.80E-06 0.43	2369 1.80E-06 0.43	1.80E-06 0.43	0.43	-	L -	0.64		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
Incg 2411 2458 0.0045 -0.23 (2458 0.0045 -0.23	2458 0.0045 -0.23	0.0045 -0.23	-0.23	\vdash		0.22		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHEKIN INCG 13
Incg 2592 2681 0.00014 0.41	2681 0.00014 0.41	2681 0.00014 0.41	0.00014 0.41	0.41	1		0.25		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
Incg 2692 2788 3.60E-20 0.44	2788 3.60E-20	2788 3.60E-20	3.60E-20	1	0.44		0.7		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
Incg 3039 3106 9.00E-07 0.45	3039 3106 9.00E-07 0.45	3106 9.00E-07 0.45	9.00E-07 0.45	0.45	\top	<u> </u>	0.51		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
Incg 3120 3191 0.00018 0.06	3191 0.00018	3191 0.00018	0.00018		90.0		0.34		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
Incg 3225 3312 1.10E-12 0.28	3312 1.10E-12 0.28	3312 1.10E-12 0.28	1.10E-12 0.28	0.28		Ϊ.	0.45		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
lncg 716 811 1.10E-21 0.16	716 811 1.10E-21	811 1.10E-21	1.10E-21		0.16		9.0		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
lncg 852 900 0.00036 -0.19	852 900 0.00036	900 0.00036	0.00036		-0.19		0.53		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
Incg 932 1021 1.60E-05 0.55	932 1021 1.60E-05	1021 1.60E-05	1.60E-05		0.55	_	0.46		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CALHERIN INCG 13
Inci B 1065 1129 3.60E-06 0.06	1065 1129 3.60E-06	1129 3.60E-06	3.60E-06	3.60E-06	90.0		0.78		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
Inci B 1178 1234 1.80E-05 -0.16	1178 1234 1.80E-05	1234 1.80E-05	1.80E-05	1.80E-05	-0.16		6.0		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
1nci B 1350 1440 3.60E-16 0.53	1350 1440 3.60E-16	1440 3.60E-16	3.60E-16	3.60E-16	0.53		90.0		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN
Inci B 1481 1547 5.40E-05 0.21	1481 1547 5.40E-05	1547 5.40E-05	5.40E-05		0.21		0.16		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN. INCI 13
1nci B 1667 1750 1.40E-06 0.26	1667 1750 1.40E-06	1750 1.40E-06	1.40E-06	1.40E-06	0.26	1	0.64		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
Inci B 181 248 5.40E-05 -0.17	181 248 5.40E-05	248 5.40E-05	5.40E-05	Ħ	-0.17	1	0.4		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN

PDB annotation	INCI 13	CELL ADHESION PROTEIN CADHERIN INCI 13	CELL ADHESION PROTEIN CADHERIN 1NCI 13	CELL ADHESION PROTEIN CADHERIN 1NCI 13	CELL ADHESION PROTEIN CADHERIN INCI 13	CELL ADHESION PROTEIN CADHERIN 1NCI 13	CELL ADHESION PROTEIN CADHERIN 1NCI 13	CELL ADHESION PROTEIN CADHERIN INCI 13	CELL ADHESION PROTEIN CADHERIN 1NCI 13	CELL ADHESION PROTEIN CADHERIN INCI 13	CELL ADHESION PROTEIN CADHERIN INCI 13	CELL ADHESION PROTEIN CADHERIN INCI 13	CELL ADHESION PROTEIN CADHERIN INCI 13	CELL ADHESION PROTEIN CADHERIN INCI 13	CELL ADHESION PROTEIN CADHERIN INCI 13	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN		
Coumpound		N-CADHERIN; INCI 3	N-CADHERIN; 1NCI 3	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3	N-CADHERIN; 1NCI 3	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3	N-CADHERIN; 1NCI 3	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN: CHAIN: A:
SeqFold score																					
PMF score		0.27	0.07	0.3	9.0	0.49		0	0.78	99.0	0.17	0.17	0.88	0.84	0.59	-1	1	0.16	. 66.0	0.88	_
Verify score		-0.5	0.24	-0.4	0.39	-0.03	0.56	-0.01	-0.24	0.52	-0.2	0.31	0.01	0.26	0.4	0.39	0.53	0.1	0.28	0.02	0.46
PSI- BLAST		0.0045	1.80E-14	3.60E-06	5.40E-07	0.0079	1.80E-19	1.80E-06	0.0045	1.10E-06	0.0013	1.40E-10	5.40E-20	7.20E-05	7.20E-06	1.10E-35	1.40E-31	1.80E-23	1.60E-58	3.60E-35	1.10E-33
End		0961	2062	2264	2371	2458	2789	2898	3003	3105	3191	3313	812	917	1022	1234	1338	1440	1547	1652	1750
Start		1912	1975	2180	2307	2412	2692	2832	2951	3041	3146	3225	716	853	932	1039	1144	1251	1351	1455	1562
Chain 10		æ	В	В	В	В	В	В	В	В	m	æ	Д	æ	ф	¥	¥	V	4	4	A
PDB ID		Inci	1 nci	Inci	Inci	Inci	Inci	Inci	Inci	1nci	Inci	Inci	Inci	lnci	Inci	Incj	Incj	Incj	Incj	Incj	Incj
SEQ S B S		608	808	608	808	608	808	808	809	809	608	809	809	809	809	809	809	809	809	809	808

																					_
PDB annotation	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL
Coumpound	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;
SeqFold score													120.25								
PMF	0.84	0.23	-	6.0	0.45	68.0	-		0.8	9.0	0.51	_			0.99		9.0	0.82	_	_	-0.03
Verify	0.2	0.08	0.31	0.25	0.02	0.23	0.25	0.32	-0.03	90.0	0.28	0.23		0.28	0.28	0.35	0.16	0.35	99.0	9.6	0.35
PSI- BLAST	7.20E-33	3.60E-22	9.00E-27	3.60E-31	1.60E-56	9.00E-31	3.60E-40	3.60E-35	9.00E-31	3.60E-40	1.30E-26	1.40E-31	3.60E-63	3.60E-63	9.00E-36	9.00E-30	1.30E-30	1.30E-30	1.80E-39	1.80E-17	5.40E-10
End	1861	354	1960	2062	2163	2264	2371	2473	2577	2683	450	2789	2897	2898	3003	3105	3196	3313	3418	3523	3621
Start	1667	173	1793	1898	1975	2079	2180	2300	2407	2488	256	2592	2691	2693	2825	2915	3039	3141	3225	3349	3473
Chain ID .	A	∢	▼	<	¥	\ ∀	<	A	V V	4	4	¥	∢	<	4	∢	4	4	4	4	4
PDB CD	Incj	Incj	Incj	Incj	Incj	Incj	1ncj	lncj	Incj	Incj	lncj	Incj	1ncj	Incj	Incj	Incj	Incj	lncj	Incj	lncj	inci
SEQ	00 00 00 00 00 00 00 00 00 00 00 00 00	809	608	608	808	809	608	809	809	808	808	608	808	608	809	608	608	808	<u></u>	808	808

PDB annotation		ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	SERINE PROTEASE FVIIA; FVIIA; BLOOD COAGULATION, SERINE	PROLEASE		SERINE PROTEASE FVIIA; BLOOD COAGULATION, SERINE	PROTEASE		SERINE PROTEASE FVIIA; FVIIA;	BLOOD COACOLATION, SEAME PROTEASE			CELL ADHESION UVOMORULIN;	CADHEKIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN;	CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL
Coumpound			N-CADHERIN; CHAIN: A;	COAGULATION FACTOR VIIA (LIGHT CHAIN); CHAIN:	L; COAGULATION FACTOR VIIA (HEAVY CHAIN);	CHAIN: H; TRIPEPTIDYL INHIBITOR: CHAIN: C;	COAGULATION FACTOR VIIA (LIGHT CHAIN); CHAIN:	L; COAGULATION FACTOR	VIIA (HEAV T CHAIN); CHAIN: H; TRIPEPTIDYL INHIBITOR: CHAIN: C:	COAGULATION FACTOR	VIIA (LIGHT CHAIN); CHAIN: L; COAGULATION FACTOR	VIIA (HEAVY CHAIN);	CHAIN: H; IMPEPILDYL INHIBITOR: CHAIN: C:	EPITHELIAL CADHERIN;	CHAIN: NULL;	EPITHELIAL CADHERIN;	CHAIN: NULL;	EPITHELIAL CADHERN; CHAIN: NULL;						
SoaFold	score																							
DME	score		0.21	0.36	96.0	0.01	1	0.88	1	0.57			-0.08			-0.15				0.13		0.27		0.45
Varific	score		-0.28	0.08	0.36	-0.13	0.39	0.35	0.48	99.0			10:0			80.0				0.33		0.24		0.04
Der	BLAST		1.10E-59	1.80E-35	7.20E-28	1.80E-25	1.10E-63	1.80E-34	3.60E-34	1.20E-21			3.60E-13			7.20E-11				7.20E-08		7.80E-20		1.30E-17
2	¥¥		248	556	299	812	917	1022	1129	3947			3976			4168				1133		1133		1238
	AA		39	390	467	573	717	827	932	3865			3904			4082				1026		1041		1145
	E A		V	¥	¥	A	4	A	A	1			L			ı								
	2 8		Incj	Incj	Incj	Incj	lncj	Incj	Incj	1qfk			lqfk			1qfk				Isuh		Isuh		1suh
	38	ÿ	608	608	608	608	608	608	809	809			809			809				809		808		809

										DDDtottori
SEQ	PDB DD	Chain	Start	End	PSI- BLAST	Verify score	PMF	SeqFold score	Coumpound	FDB alliciation
ÿ										ADHESION
809	1suh		1171	1238	7.20E-08	-0.03	0.4		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMOKULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
809	1 suh		1249	1334	1.30E-10	0.54	0.62		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
608	1suh		1350	1444	1.80E-21	0.3	0.25		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
608	Isuh		1455	1549	1.30E-08	0.45	0.63		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
608	1suh		1455	1551	5.40E-07	0.14	0.11		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMUKULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
809	Isuh		155	226	2.60E-07	0.35	0.07		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UYOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
809	1 suh		1570	1650	1.00E-12	0.41	0.31		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
809	1suh		1589	1656	1.40E-08	0.2	60:0		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UYOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
809	Isuh		1667	1754	1.30E-12	0.07	99.0		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UYOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	1suh		1690	1754	3.60E-06	0.06	0.66	_	EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UYOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
809	1suh		1777	1867	1.30E-14	-0.06	0.48		EPITHELJAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
809	Isuh		1780	1868	3.60E-05	-0.24	0.39		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	1suh		182	252	5.40E-07	0.41	0.13		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	1suh		1898	1964	1.60E-05	-0.1	0.18		EPITHELIAL CADHERIN;	CELL ADHESION UVOMORULIN;

PDB annotation	CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UYOMORULIN; CADHERIN, CALCTUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UYOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCTUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UYOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
PD	CADHERIN, CA ADHESION	CELL ADHESIO CADHERIN, CA ADHESION	CELL ADHESIC CADHERIN, CA ADHESION	CELL ADHESIC CADHERIN, CA ADHESION	CELL ADHESIC CADHERIN, CA ADHESION	CELL ADHESIC CADHERIN, CA ADHESION	CELL ADHESIC CADHERIN, CA ADHESION	CELL ADHESIC CADHERIN, CA ADHESION	CELL ADHESIC CADHERIN, CA ADHESION	CELL ADHESIC CADHERIN, CA ADHESION	CELL ADHESIC CADHERIN, CA	CELL ADHESIC CADHERIN, CA	CELL ADHESIC CADHERIN, CA	CELL ADHESIC CADHERIN, CA
Coumpound	CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;
SeqFold score														
PMF score		1202.08	0.09	0.12	0.57	0.46	0.77	0.94	0.58	90.00	0.03	0.98	0.12	66:0
Verify score		0.17	0.42	0.05	-0.23	-0.36	0.44	0.4	0.03	-0.37	-0.2	0.16	0.02	0.51
PSI- BLAST		7.20E-18	2.60E-15	3.60E-07	1.30E-14	3.60E-10	3.90E-20	1.10E-08	3.90E-05	0.00011	3.60E-12	1.30E-13	2.60E-10	1.40E-23
End		2066	2163	2167	2268	2268	2375	2375	2475	2477	2581	2581	2681	2793
Start		1975	2084	2104	2178	2178 ·	2280	2306	2395	2414	2488	2489	2591	2692
Chain TD														
PDB ID		1suh	1suh	1suh	1suh	1suh	Isuh	1suh	1suh	1suh	1suh	1suh	1suh	Isuh
SEQ	Ö	808	809	809	809	809	809	808	808	808	608	808	808	808

PDB annotation														4; CELL ADHESION UVOMORULIN;
Coumpound	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN; NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL:	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN;
SeqFold score														
PMF	0.87	0.53	0.93	0.86	68.0	0.55	0.25	0.82	0.98	0.94	0.65	0.24	0.3	0.19
Verify	0.43	-0.01	0.37	-0.07	0.36	0.33	0.43	9.0	0.58	9.0	0.56	0.08	-0.21	-0.03
PSI- BLAST	2.60E-21	1.40E-08	1.20E-14	1.80E-05	2.60E-21	3.60E-06	3.90E-12	1.30E-13	3.90E-21	5.40E-05	6.50E-11	0.0001	9.00E-06	1.30E-15
End	2902	2902	3007	3007	3109	3109	3213	3317	3422	3422	3524	454	454	260
Start	2806	2831	2914	2941	3026	3045	3120	3225	3330	3355	3435	363	401	467
Chain														
PDB	1suh	Isuh	1suh	Isuh	1 suh	Isuh	1suh	1suh .	1suh	lsuh	Isuh	1suh	1suh	Isuh
SEQ	809 809	809	809	809	808	809	809	809	808	808	808	608	809	808

PDB annotation		ADHESION	CELL ADHESION UYOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	PLASMINOGEN ACTIVATION	BLOOD COAGULATION FACTOR STUART FACTOR, BLOOD COAGULATION FACTOR, SERINE PROTEINASE, EPIDERMAL 2 GROWTH FACTOR LIKE DOMAIN		CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN				
Coumpound			EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	T-PLASMINOGEN ACTIVATOR F1-G; 1TPG 7 CHAIN: NULL; 1TPG 8	BLOOD COAGULATION FACTOR XA; CHAIN: L, C;	LECTIN (AGGLUTININ) WHEAT GERM AGGLUTININ (ISOLECTIN 2) 9WGA 3	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;
SeqFold	score												
PMF	score		0.57	0.95	0.72	0.47	0.57	0.18	0.59	0.1	-0.2	0.25	0.16
Verify	score		0.39	-0.21	0.4	0.27	0.37	0.2	0.21	-0.26	0.02	-0.26	0.35
PSI-	BLAST		3.90E-05	7.20E-25	1.00E-17	1.30E-07	5.20E-13	9.00E-07	1.30E-18	7.20E-11	5.40E-12	3.60E-24	1.80E-17
End	*		658	816	921	921	1026	1026	3940	3994	4234	370	145
Start	¥¥		290	716	826	854	930	959	3863	3904	4069	190	23
Chain	A									<u>-1</u>	4	∢	٧
PDB	А		1suh	1suh	1suh	1suh	1suh	1suh	ltpg	Ixka	9wga	ledh	ledh
SEO	АŞ		608	808	808	808	808	808	808	809	809	811	811

PDB annotation	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELLIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL AHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELLAL CADHERIN DOMAINS 1 AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCI 13	CELL ADHESION PROTEIN CADHERIN INCI 13	CELL ADHESION PROTEIN CADHERIN INCI 13	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL
Coumpound	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN: CHAIN: A;	N-CADHERIN; CHAIN: A;
SeqFold score			2	124.11											
PMF	0.82	-	-		0.03	0.63	_	0.12	0.62	1	96.0	0.59	0.58	0.99	_
Verify score	0.21	0.21	0.61		0.07	0.34	0.16	-0.15	0.44	0.35	0.21	0.31	0.31	0.24	0.29
PSI- BLAST	1.805-30	7.20E-26	1.80E-51	1.80E-51	0.00018	1.80E-06	1.60E-19	0.00036	3.60E-07	5.40E-19	1.10E-26	3.60E-20	1.80E-31	7.20E-28	1.80E-55
End	474	584	237	254	238	473	143	238	474	145	370	145	474	999	238
Start	296	410	20	20	188	408	48	189	409	48	160	24	27.1	402	49
Chain	V	V	4	∢				m	æ	æ	∢	4	∢	<	٧
PDB ID	ledh	ledh	1edh	1edh	Incg	Incg	Incg	lnci	Inci	1nci	lncj	Incj	Incj	Incj	Incj
SEQ EQ	811 118	811	811	811	811	811	811	811	811	811	811	811	811	811	811

													· · · · · · · · · · · · · · · · · · ·
ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN: CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGERDNA) COMPLEX (ZINC FINGERDNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGERDNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
	N-CADHERIN; CHAIN: A;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	QGSR ZINC FINGER	PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE: CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE: CHAIN: B. C;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;
	125.8												
		0.18	0.23	0.01	0.46	0.89	-	-	0.1			0.92	-
		0.22	-0.08	0.36	0.3	-0.15	0.65	0.58	0.23	0.61		0.68	0.34
	1.80E-55	1.20E-16	0.0041	5.20E-12	1.30E-17	1.60E-08	9.00E-23	2.60E-29	1.00E-08	2.60E-35		3.60E-27	7.205-49
	253	258	237	371	478	478	149	149	584	142		142	198
	49	162	061	268	383	410	48	49	490	62		69	117
	¥.									₹		4	၁
1	Incj	1suh	lsuh	1suh	1suh	1suh	Isuh	1suh	1suh	lalh		lalh	Ime y
ö	811	811	811	811	811	811	811	811	811	813		813	813
	NO: ADHESION PROTEIN	1ncj A 49 253 1.80E-55 125.8 N-CADHERIN; CHAIN: A;	1ncj A 49 253 1.80E-55 125.8 N-CADHERIN; CHAIN: A; 15uh 162 258 1.20E-16 0.22 0.18 EPITHELIAL CADHERIN; CHAIN: NULL;	1ncj A 49 253 1.80E-55 125.8 N-CADHERIN; CHAIN: A; 15uh 162 258 1.20E-16 0.22 0.18 EPITHELIAL CADHERIN; CHAIN: NULL; 15uh 190 237 0.0041 -0.08 0.23 EPITHELIAL CADHERIN; CHAIN: NULL; CHAIN	1ncj A 49 253 1.80E-55 125.8 N-CADHERIN; CHAIN: A; 15uh 162 258 1.20E-16 0.22 0.18 EPITHELIAL CADHERIN; CHAIN: Hsuh 190 237 0.0041 -0.08 0.23 EPITHELIAL CADHERIN; CHAIN: NULL; 15uh 268 371 5.20E-12 0.36 0.01 EPITHELIAL CADHERIN; CHAIN: NULL; CHAIN: NUL	1 1 1 1 2 253 1.30E-15 0.18 125.8 N-CADHERIN; CHAIN: A; 15 1.30E-16 0.22 0.18 EPITHELIAL CADHERIN; CHAIN: NULL; 15 1.30E-17 0.36 0.01 EPITHELIAL CADHERIN; CHAIN: NULL; CH	1 1 1 1 2 253 1.80E-55 1.25.8 N-CADHERIN; CHAIN: A; 1.5 1.20E-16 0.22 0.18 EPITHELIAL CADHERIN; CHAIN: NULL; CH	1 1 1 1 2 253 1.80E-55 1.25.8 N-CADHERIN; CHAIN: A; 1.5 1.5 1.20E-16 0.22 0.18 EPITHELIAL CADHERIN; CHAIN: NULL; 1.5	1ncj A 49 253 1.80E-55 125.8 N-CADHERIN; CHAIN: A; Chain	11cg A 49 253 1.80E-55 1.25.8 N-CADHERIN; CHAIN: A; 15th 162 258 1.20E-16 0.22 0.18 EPITHELIAL CADHERIN; CHAIN: NULL; CHAIN:	1ncj A 49 233 1.80E-55 1.25.8 N-CADHERIN; CHAIN: A; 1.80E-55 1.20E-16 0.22 0.18 EPITHELIAL CADHERIN; CHAIN: NULL; 1suh 190 237 0.0041 -0.08 -0.23 CHAIN: NULL; CH	1ncj A 49 253 1.80E-55 125.8 N-CADHERIN; CHAIN; A; 151.8 N-CADHERIN; CHAIN; CHAIN; NULL; 152.8 1.20E-16 0.22 0.18 EPITHELIAL CADHERIN; CHAIN; NULL; CHAIN; CHAIN	Incj A 49 253 1.80E-55 125.8 N-CADHERIN; CHAIN: A; 1.80E-56 1.20E-16 0.22 0.18 EPITHELIAL CADHERIN; CHAIN: A; 1.20E-17 0.0041 -0.08 0.23 EPITHELIAL CADHERIN; CHAIN: NULL; CHAIN: NU

PDB annotation	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGERODA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA).	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA FINERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	·
Coumpound	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	TRANSCRIPTION REGULATION YEAST TRANSCRIPTION FACTOR ADR I (RESIDUES 130 - 159) IPAA 3 (PAPA - CARBOXY TERMINAL ZINC FINGER
SeqFold				111.37				
PMF	-		-		1	1	1	0.34
Verify score	0.51	0.32	0.34		0.27	0.63	0.39	0.13
PSI- BLAST	5.40E-50	1.80E-49	1.30E-49	1.30E-49	1.30E-49	9.00E-43	1.10E-48	9.00E-05
End	226	254	282		310	142	170	312
Start AA	145	173	201	201	229	89	68	286
Chain	U	U	U	U	υ	ပ	U	
PDB	line y	y me	y Tae	y y	Jme y	Jme y	Jmc y	l paa
SEQ		813	813	813	813	813	813	813

PDB annotation		ZINC FINGER TRANSCRIPTION FACTOR SPI; ZINC FINGER, TRANSCRIPTION ACTIVATION, SPI	COMPLEX (TRANSCRIPTION REGULATIONDNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POLYMERASE JII, 2 TRANSCRIPTION INITATION, ZNC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INITATION, ZNC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INITATION, ZNC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YYI, ZINC 2 RECOGNITION, 3 COMPLEX (TRANSCRIPTION, 3 COMPLEX (TRANSCRIPTION REGULATION/DA)
Coumpound	DOMAIN) MUTANT WITH IPAA 4 PRO 131 REPLACED BY ALA, PRO 133 REPLACED BY ALA, CYS 140 IPAA 5 REPLACED BY ALA (P131A,P133A,C140A) (NMR, 10 STRUCTURES) IPAA 6	SP1F2; CHAIN: NULL;	TFIIIA; CHAIN: A, D; SS RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	TFIIIA; CHAIN: A, D; SS RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;
SeqFold score				111.05			
PMF score		0.35	0.98		-	p===	-
Verify score		0.27	0.17		0.36	0.38	0.26
PSI- BLAST		5.40E-07	1.805-38	2.60E-61	9.00E-36	7.20E-37	2.60E-49
End		312	270	310	312	235	254
Start	·	286	118	145	174	8	144
Chain ID			∢	∢	∢	<	υ
PDB ID		1sp2	146	1tf6	1116	1116	lubd
S B S		813	813	813	813	813	813

		<u> </u>				<u> </u>
PDB annotation	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG I; TRANSCRIPTION INTIATION, INTIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INTIATION, INTIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG I; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YYI, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY!, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX CTRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX
Coumpound	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI, CHAIN: C; ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;	YY I; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YY I; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INTIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;
SeqFold score		105.14				
PMF	1	·			0.94	86:0
Verify score	0.25		0.03	0.37	0.49	0.34
PSI- BLAST	3.60E-35	1.30E-50	7.80E-50	1.30E-50	2.60E-43	1.80E-33
End		283	283	310	170	170
Start AA	153	. 221	178	199	99	69
Chain ID	ပ	ပ	ပ	ပ	ပ	U
PDB	lubd	lubd	1ubd	lubd	lubd	Iubd
SEQ SO BEQ	813	813	813	813	813	813

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PDB annotation	DOMAIN, BETA-BARREL, MIXED ALPHA-BETA, HEXAMER, 2 DIMER	SCAFFOLD PROTEIN SCAFFOLD PROTEIN, PP2A, PHOSPHORYLATION, HEAT REPEAT	TOXIN BINDING PROTEIN TWO DOMAINS: BETA PROPELLER AND ALPHA/BETA FOLD	TRANSCRIPTION INHIBITOR BETA- PROPELLER	TRANSCRIPTION INHIBITOR BETA- PROPELLER	COMPLEX (GTP-	TRANSDUCIN BETA SUBUNIT:	GAMMAI, TRANSDUCIN GAMMA	SUBUNIT; COMPLEX (GTP-	BINDING/I KANSDUCEK), G PROTEIN, HETEROTRIMER 2 SIGNAL	TRANSDUCTION	COMPLEX (GTP-	TRANSDUCIN BETA SUBUNIT;	GAMMAI, TRANSDUCIN GAMMA	SUBUNIT: COMPLEX (GTP-	BINDING/TRANSDUCER), G PROTEIN,	TRANSDUCTION	NUCLEAR TRANSPORT PROTEIN	TRANSPORT PROTEIN COMPLEX	ARMADILLO REPEAT ARMADILLO	REPEAT, BETA-CATENIN,	CYTOSKELETON	OCADI TO GIGDS AGICD AT 100 CO	COMPLEX (HSP24/HSP70) HSP70, GRPE, MOLECULAR CHAPERONE, NUCLEOTIDE EXCHANGE 2 FACTOR.
Coumpound	METHYLTRANSFERASE; CHAIN: 1, 2, 3, 4, 5, 6;	PROTEIN PHOSPHATASE PP2A; CHAIN: A, B;	TOLB PROTEIN; CHAIN: A;	TRANSCRIPTIONAL REPRESSOR TUP1; CHAIN: A, B, C;	TRANSCRIPTIONAL REPRESSOR TUP1; CHAIN: A, B, C;	GT-ALPHA/GI-ALPHA	BETA: CHAIN: A; GI-	GAMMA; CHAIN: G;				GT-ALPHA/GI-ALPHA CHIMERA: CHAIN: A: GT.	BETA; CHAIN: B; GT-	GAMMA: CHAIN: G;				KARYOPHERIN BETA2;	CITAIN: D. NAW, CHAIN: C.	BETA-CATENIN; CHAIN:	NULL;		MICH EOTHER EVOLANCE	NOCLEO ILLE EXCHANGE FACTOR GRPE; CHAIN: A, B; MOLECULAR CHAPERONE
SeqFold score										•													70.02	78.03
PMF score		0.98	0.03	-	1	-						•						0.07		0.23				
Verify score		0.32	0.16	0.12	0.28	0.18						0.63						-0.04		-0.02				
PSI- BLAST		1.20E-07	0.0036	3.60E-60	3.60E-68	3.60E-60						3.60E-73						5.20E-05		2.60E-15			5 20E-41	3.202-1
End		520	1078	1099	1147	1096						1144						653		653			222	
Start AA		235	821	796	831	758						823						198		345			51	;
Chain		<	∢	V.	¥	В						ф						ф					4	:
PDB TD		163u	1crz	lerj	lerj	1got						1got						145k		3bct			1dke	
SEQ NO H		817	817	817	817	817						817						817		817			818	

				_	·					
PDB annotation	COLLED-COIL, COMPLEX (HSP24/HSP70)	COMPLEX (HSP24/HSP70) HSP70, GRPE, MOLECULAR CHAPERONE, NUCLEOTIDE EXCHANGE 2 FACTOR, COILED-COIL, COMPLEX (HSP24/HSP70)	COMPLEX (HSP24/HSP70) HSP70, GRPE, MOLECULAR CHAPERONE, NUCLEOTIDE EXCHANGE 2 FACTOR, COILED-COIL, COMPLEX (HSP24/HSP70)		COMPLEX (DNA-BINDING PROTEIN/DNA) MYN PROTEIN; MAX, DNA BINDING, BASIC-HELIX-LOOP- HELIX-LEUCINE ZIPPER, 2 TRANSCRIPTION FACTOR, COMPLEX (DNA-BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) MYN PROTEIN; MAX, DNA BINDING, BASIC-HELIX-LOOP- HELIX-LEUGINE ZIPPER, 2 TRANSCRIPTION FACTOR, COMPLEX (DNA-BINDING PROTEIN/DNA)	COMPLEX (TRANSCRIPTION FACTOR MAXIDNA) TRANSCRIPTIONAL REGULATION, DNA BINDING, COMPLEX 2 (TRANSCRIPTION FACTOR MAXIDNA)	COMPLEX (TRANSCRIPTION FACTOR MAX/DNA) TRANSCRIPTIONAL REGULATION, DNA BINDING, COMPLEX 2 (TRANSCRIPTION FACTOR MAX/DNA)	SCAFFOLD PROTEIN SCAFFOLD PROTEIN, PP2A, PHOSPHORYLATION, HEAT REPEAT	SMALL GTPASE KARYOPHERIN BETA, P95 SMALL GTPASE, NUCLEAR
Coumpound .	DNAK; CHAIN: D;	NUCLEOTIDE EXCHANGE FACTOR GRPE; CHAIN: A, B; MOLECULAR CHAPERONE DNAK; CHAIN: D;	NUCLEOTIDE EXCHANGE FACTOR GRPE; CHAIN: A. B; MOLECULAR CHAPERONE DNAK; CḤAIN: D;		MAX PROTEIN; CHAIN: A, C; DNA; CHAIN: B, D;	MAX PROTEIN; CHAIN: A, C; DNA; CHAIN; B, D;	TRANSCRIPTION FACTOR MAX; CHAIN: A. B. DNA (5'- D(*CP*AP*CP*CP*AP*CP*GP *TP*GP*GP*T)-3', CHAIN: C, D;	TRANSCRIPTION FACTOR MAX; CHAIN: A, B; DNA (5:- D(*CP*AP*CP*CP*AP*CP*GP *TP*GP*GP*Ty-3', CHAIN: C, D;	PROTEIN PHOSPHATASE PP2A; CHAIN: A, B;	RAN; CHAIN: A, C; IMPORTIN BETA SUBUNIT;
SeqFold score										
PMF		0.83	0.42		0.03	0:00	0	0.28		0.82
Verify score		-0.15	-0.29		-0.73	8.0-	-0.58	-0.27	0.55	0.25
PSI- BLAST		1.80E-20	5.20E-41		3.60E-14	2.60E-13	1.60E-14	1.80E-14	6.50E-08	0.0026
End AA		221	220		203	219	203	203	373	235
Start AA		53	19		135	140	131	132	44	131
Chain ID		∀	4		₹	₹	∢	м	V	В
PDB		ldkg	ldkg		lan2	lan2	1hlo	1hlo	1b3u	1jbr
SEQ SEQ		818	818		820	820	820	820	824	824

	_			 					_						_		_						
PDB annotation	TRANSPORT RECEPTOR	SMALL GTPASE KARYOPHERIN BETA, P95 SMALL GTPASE, NUCLEAR TRANSPORT RECEPTOR	ARMADILLO REPEAT ARMADILLO REPEAT, BETA-CATENIN, CYTOSKELETON	LYASE ACC SYNTHASE, S- ADENOSYL-L-METHIONINE ETHYLENE BIOSYNTHESIS	RIFAMYCIN BIOSYNTHESIS (RIFD GENE) AHBA SYNTHASE; RIFAMYCIN BIOSYNTHESIS (RIFD GENE)	TRANSFERASE TRANSFERASE, METABOLIC ROLE, PYRIDOXAL 5'- PHOSPHATE	TO A UNION A COLUMN A	AMINOTRANSFERASE, PYRIDOXAL ENZYME	TRANSFERASE AONS, 8-AMINO-7-	KETOPELARGONATE SYNTHASE; PLP-DEPENDENT ACYL-COA	SYNTHASE, BIOTIN BIOSYNTHESIS,	8-2 AMINO-7-OXONANOATE	SYNTHASE, 8-AMINO-/- KETOPEL ARGONATE 3 SYNTHASE	TRANSFERASE	LYASE ALPHA/BETA FOLD	TRANSFERASE SHMT; HYDROXYMETHYL TRANSFERASE	CARBON METABOLISM	METHIONINE BIOSYNTHESIS BETA	CYSTATHIONASE; PLP-DEPENDENT	BIOSYNTHESIS C-S BETA 21.YASE	LYASE CGS; LYASE, LLP-DEPENDENT	ENZYMES, METHIONINE BIOSYNTHESIS	TRANSFERASE AMINOTRANSFERASE FOLD, LARGE PLP-BINDING DOMAIN,
Coumpound	CHAIN: B, D;	RAN; CHAIN: A, C; IMPORTIN BETA SUBUNIT; CHAIN: B, D;	BETA-CATENIN; CHAIN: NULL;	1-AMINOCYCLOPROPANE-1- CARBOXYLATE SYNTHASE; CHAIN: A, B;	3-AMINO-5- HYDROXYBENZOIC ACID SYNTHASE; CHAIN: A;	SERINE HYDROXYMETHYLTRANSF FRASE: CHAIN: A:	ACDADITATES	ASPARIALE AMINOTRANSFERASE; CHAIN: A. B:	8-AMINO-7-OXONANOATE	SYNTHASE; CHAIN: A;					CSDB PROTEIN; CHAIN: A;	SERINE HYDROXYMETHYLTRANSF	ERASE; CHAIN: A, B;	CYSTATHIONINE BETA-	LYASE; CHAIN: A, B;		CYSTATHIONINE GAMMA-	SYNTHASE; CHAIN: A, B, C, D;	MALY PROTEIN; CHAIN: A, B;
SeqFold score									52.33							•		•					
PMF		0.47	0.99	0.22	-0.13	0		1202.08							_	-1.41		61.0-			-0.03		0.51
Verify score		90:0	0.41	0.41	0.12	0.25	000	67:0							0.51	0.04		0.1			0.2		-0.06
PSI- BLAST		6.50E-06	6.50E-14	5.40E-14	3.60E-37	5.40E-50	1 900 60	1.005-32	7.20E-06						9.00E-57	3.60E-51		3.60E-16			5.40E-34		1.80E-09
End		303	404	294	317	312	300	Sec.	317						311	312		314			313		259
Start AA		201	154	07	26	-	,	<u> </u>	-						10	-		83			22		135
Chain ID		В		٧	٧	¥	<	¢	¥		•				∀	∢		¥			4		¥
PDB UD		1ibr	3bct	1b8g	159h	1bj4	I Pilin	<u> </u>	1650						1con	100		Icli			1cs1		1d2f
SEQ BÖ		824	824	826	826	826	32	3	826						826	826		826			826		826

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PDB annotation	SMALL C-2 TERMINAL DOMAIN, OPEN ALPHA-BETA STRUCTURE.	TRANSFERASE SHMT, SERINE METHYLASE; ALPHA PLP ASPARTATE, AMINO TRANSFERASE, (AAT)-LIKE FOLD	TRANSFERASE PLP-DEPENDENT ENZYMES, IRON-SULFUR-CLUSTER SYNTHESIS, C-S 2 BETA LYASE	TRANSFERASE SHMT; SERINE- GLYCINE CONVERSION, PYRIDOXAL S-PHOSPHATE, 2 TETRAHYDROFOLATE, ASYMMETRIC DIMER	LYASE METHIONINE BIOSYNTHESIS, PYRIDOXAL 5'-PHOSPHATE, GAMMA- 2 FAMILY, LYASE		LYASE LYASE, PLP-DEPENDENT ENZYME, PYRIDOXAL PHOSPHATE	LIGASE EGAP; UBCH7; BILOBAL STRUCTURE, ELONGATED SHAPE, E3 UBIQUITIN LIGASE, E2 2 UBIQUITIN CONJUGATING ENZYME	LIGASE E6AP; UBCH7; BILOBAL STRUCTURE, ELONGATED SHAPE, E3 UBIQUITIN LIGASE, E2 2 UBIQUITIN CONJUGATING ENZYME	SH3 PROTOTYPE WWPROTOTYPE, PROTEIN DESIGN	SH3 PROTOTYPE WWPROTOTYPE, PROTEIN DESIGN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA)
Coumpound		SERINE HYDROXYMETHYLTRANSF ERASE; CHAIN: A, B, C, D;	AMINOTRANSFERASE; CHAIN: A, B;	SERINE HYDROXYMETHYLTRANSF ERASE; CHAIN: A, B, C, D;	CYSTATHIONINE GAMMA- SYNTHASE; CHAIN: A, B, C, D, E, F, G, H;	LYASE(CARBON-CARBON) TYROSINE PHENOL-LYASE (E.C.4.1.99.2) 1TPL 3	TYROSINE PHENOL-LYASE; CHAIN: A, B;	UBIQUITIN-PROTEIN LIGASE E3A; CHAIN: A, B, C; UBIQUITIN CONJUGATING ENZYME E2; CHAIN: D;	UBIQUITIN-PROTEIN LIGASE E3A; CHAIN: A, B, C; UBIQUITIN CONJUGATING ENZYME E2; CHAIN: D;	WWPROTOTYPE; CHAIN: A;	WWPROTOTYPE; CHAIN: A;	OGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE DENDING STEE. CHAIN: D. C.	QGSR ZINC FINGER
SeqFold score								217.98					
PMF		0.17	-	-1.41	-0.06	-0.03	-0.06		-	0.29	0.35	0.24	-
Verify score		-0.07	0.62	0.09	0.19	0.21	0.02		0.38	90.0	0.01	-0.37	0.04
PSI- BLAST		9.00E-54	3.60E-58	5.40E-50	7.20E-21	3.60E-09	1.30E-08	1.10E-45	1.10E-45	2.60E-11	3.90E-12	3.60E-26	2.60E-39
End		314	317	312	311	311	311	1566	1565	841	1019	259	315
Start		œ	32	-	20	101	101	1213	1239	808	985	179	235
Chain		4	4	⋖	⋖	∢	4	A	∢	4	<	A	4
PDB ID		gp1	Zgel Zgel	leji	lqgn	lţ.	2tpl	1042	1c4z	은 등 등	9 E	lalh	lalh
SEQ PO PO PO PO PO PO PO PO PO PO PO PO PO		826	826	826	826	826	826	830	830	830	830	832	832

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PDB annotation	COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CXYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CYYSTAL STRUCTURE, COMPLEX (ZINC FINGER DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CYYSTAL STRUCTURE, COMPLEX	COMPLEX (ZINC FINGER/DNA) ZINC
Coumpound	PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B. C;	QGSR ZINC FINGER PEPTIDE: CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E;
SeqFold score										
PMF			0.99	0.98	0.07	0.52	_		-	_
Verify score		0.48	0.41	0.27	-0.35	-0.16	0.1	0.23	0.55	0.49
PSI- BLAST		9.10E-38	1.40E-30	3.90E-39	3.60E-40	1.10E-43	3.60E-45	1.60E-46	1.40E-47	1.30E-48
End		344	879	829	231		287	315	343	371
Start AA		263	598	598	150	178		234	262	290
Chain ID		¥	A	Ą	ပ	ပ	ပ	υ	ن	S
PDB ID		lalh	lalh	laih	Ime y	lme y	1me y	lme y	Ime y	Ime
SEQ ID NO:		832	832	832	832	832	832		832	832

PDB annotation	FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA FINGER, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGENDNA) ZINC
Coumpound	CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, B; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, B; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E, CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A. B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E;
SeqFold score									
PMF score		-	1	-	-		86.0	-	-
Verify score		0.62	0.34	0.27	0.42	0.16	-0.22	0.1	0.43
PSI- BLAST		1.80E-49	1.40E-49	3.60E-50	5.40E-50	7.20E-50	6.50E-33	1.10E-49	7.20E-48
End		399	427	455	483	511	538	539	594
Start AA		318	346	374	402	430	430	458	514
Chain ID		U	ပ	ပ	υ	ပ	ပ	ပ	ပ
PDB	^	Jme y	jme y	y y	Jmc y	Jme y	Jme y	Jme y	l Be
SEQ DO:		832	832	832	832	832	832	832	832

PDB annotation	INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA	INTERACTION, PROTEIN DESIGN, 2	CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC	FINGER, PROTEIN-DNA	INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRICTIRE, COMPLEX	(ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC	FINGER, PROTEIN-DNA	INTERACTION, PROTEIN DESIGN, 2	CRYSTAL STRUCTURE, COMPLEX	COMPLEX (ZINC FINGER/DNA) ZINC	FINGER, PROTEIN-DNA	INTERACTION, PROTEIN DESIGN, 2	CRYSTAL STRUCTURE, COMPLEX	(ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC	FINGER, PROTEIN-DNA	INTERACTION, PROTEIN DESIGN, 2	CRYSTAL STRUCTURE, COMPLEX	COMPLEX (TRANSCRIPTION	REGULATION/DNA) TFIIIA: 5S GENE:	NMR. TFIIIA. PROTEIN. DNA.	TRANSCRIPTION FACTOR, 55 RNA 2	GENE, DNA BINDING PROTEIN, ZINC	FINGER, COMPLEX 3	(TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA) TFIIIA; 5S GENE;	NMR, TFIIIA, PROTEIN, DNA,	IKANSCRIPTION FACTOR, 5S RNA 2	GENE, DNA BINDING PROTEIN, ZINC FINGER, COMPLEX 3
Coumpound	PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;		DNA; CHAIN: A, B, D, E;	CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;		DNA; CHAIN: A, B, D, E;	CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;		DNA: CHAIN: A B D E.	CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;			DNA; CHAIN: A, B, D, E;	CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;		TRANSCRIPTION FACTOR	IIIA: CHAIN: A: 5S RNA	GENE: CHAIN: E. F.					TRANSCRIPTION FACTOR	IIIA; CHAIN: A; SS RNA	GENE; CHAIN: E, F;		
SeqFold score					101.87	*																											
PMF		_							-			-	-	1				0.94				0.99							0				
Verify score	-	69.0							12.0			,	0.39	ì				90.0				0.15							-0.18				
PSI- BLAST		1.80E-50			1.80E-50				3.60E-50				1.80E-49	1				1.30E-12				1.40E-20	•						3.60E-18				
End		622			623				650				8/9					371				511							999				
Start		541			541				695				597					344				431				_			487				
Chain		ပ			ပ				၁				O					_ල				A							₹.				
PDB ID		Jac V			Ime	>			Ime	>			Ime	>				Ime	>		-	143							<u> </u>	_			
SEQ NO:		832			832				832	•			832					832				832							832				

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PDB annotation	(TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION)	REGULATION/DNA), RNA	POLYMERASE III, 2 TRANSCRIPTION INITIATION ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION	REGULATION/DNA) COMPLEX	(IKANSCKIPTION	RECOLATION/DNA), KNA POLYMERASE III. 2 TRANSCRIPTION	INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION	REGULATION/DNA) COMPLEX	(IKANSCKIPTION	REGULATION/DNA), RNA	POLYMERASE III, 2 TRANSCRIPTION	ON THE TOP AND THE PROTEIN	COMPLEX (I RANSCRIP I JON	(TEGULATION/DINA) COMPLEX	(IKANSCRIPTION	REGULATION/DNA), KNA	POLYMERASE III, 2 TRANSCRIPTION	INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION	REGULATION/DNA) COMPLEX	(IKANSCKIPIION	REGULATION/DIAA), MAA	NITIATION ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION	REGULATION/DNA) COMPLEX	(TRANSCRIPTION	REGULATION/DNA), RNA	POLYMERASE III, 2 TRANSCRIPTION	OCH TEXT OF ANICON TROPERS	REGIT ATTON/DNA) COMPLEX	(TRANSCRIPTION	REGULATION/DNA), RNA
Coumpound		TFIJIA; CHAIN: A, D; SS RIBOSOMAL RNA GENE; CHAIN: B, C, F, F.	CHAIN, B, C, L, 1,		TFIIIA; CHAIN: A, D; 5S	RIBOSOMAL RNA GENE;	CHAIN: B, C, E, F;			TFIIIA; CHAIN: A, D; 5S	RIBOSOMAL RNA GENE;	CHAIN: B, C, E, F;			C	PERIOR CHAIN: A, U; 35	KIBUSUMAL KNA GENE;	CHAIN: B, C, E, F;				TFIIIA; CHAIN: A, D; 5S	RIBOSOMAL RNA GENE;	CHAIN: B. C. E. F;	,		TFIIIA; CHAIN: A, D; 5S	RIBOSOMAL RNA GENE;	CHAIN: B, C, E, F;	•		TEHLY CHANG A P. 65	PIBOSOMAI BNA GENE	CHAIN: B. C. E. F.	
SeqFold score					105.86															•															
PMF		0.84								_					;	-1.41						-1.41					0.89					700	0.0		
Verify score		0.09								0.15		_			3	0.32						0.16					-0.05			_		200	5		
PSI- BLAST		1.80E-36			1.30E-66			-	-	1.60E-36					20 200	3.00E-37						1.30E-36					1.60E-36					1 900 35	1.00E-30		
End		352			431					408						404						575					603					631	3		
Start		207			262					263					9.5	213						431					459					107	è		
Chain		4			Ą					4						<						<					4					•	ζ		
PDB ID		941			1116					146					2	OILI						11£6					1116					1+6%	2		
SEQ B B		832			832					832					000	937						832					832					833	1		

. PDB annotation	POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA REGULATION/DNA), RNA	NITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1;	INTIATOR ELEMENT, YY1, ZINC 2	FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX	(TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGIT ATTOMONA) VING-YANG I	TRANSCRIPTION INITIATION,	INITIATOR ELEMENT, YY1, ZINC 2	RECOGNITION, 3 COMPLEX	(TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA) YING-YANG 1;	TRANSCRIPTION INITIATION,	INITIATOR ELEMENT, YYI, ZINC 2	RECOGNITION 3 COMPLEX	(TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION	TRAUSCRIPTION INITIATION	INITIATOR ELEMENT, YY1, ZINC 2	FINGER PROTEIN, DNA-PROTEIN	RECOGNITION, 3 COMPLEX	(TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION	TRANSCRIPTION INITIATION	INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN
Coumpound		TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;		YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS PS	CHAIN: A, B;			YY1; CHAIN: C; ADENO-	INITIATOR ELEMENT DNA;	CHAIN: A, B;			YYI; CHAIN: C; ADENO-	ASSOCIATED VIRUS P5	MITIATOR ELEMENT DNA;	CHAIN: A, B;			YYI; CHAIN: C; ADENO-	ASSOCIATED VIROS PS	CHAIN: A. B.				YYI; CHAIN: C; ADENO-	INITIATOR ELEMENT DNA:	CHAIN: A, B;
SeqFold score																											
PMF		-		0.18				96'0					69.0						96'0						_		
Verify		0.3		-0.36				-0.38					-0.37						-0.07						-0.01		,
PSI- BLAST		7.20E-35		9.00E-29				3.60E-31					1.30E-34						2.60E-48						3.60E-32		
End		099		259				287					315						371						343		
Start		515		158				186					188						232						242		
Chain 10		K		ပ				၁					ပ						ပ						ပ		
PDB D		1166		Iubd				1ubd	,				Inbd						pqn						pgn		
SEQ NO.		832		832				832					832						832						832	•	

PDB annotation	RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION,	INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN RECOGNITION, 3 COMPLEX	COMPLEX (TRANSCRIPTION	REGULATION/DNA) YING-YANG I; TRANSCRIPTION INITIATION.	INITIATOR ELEMENT, YYI, ZINC 2	FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX	(IRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1;	TRANSCRIPTION INITIATION,	INITIATOR ELEMENT, YY1, ZINC 2	FINGER PROTEIN, DNA-PROTEIN	(TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA) YING-YANG 1;	TRANSCRIPTION INITIATION,	INITIATOR ELEMENT, YY1, ZINC 2	FINGER PROTEIN, DNA-PROTEIN	(TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA) YING-YANG 1;	IKANSCRIPTION INITIATION,	FINGER PROTEIN DNA-PROTEIN	RECOGNITION 3 COMPLEX	(TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA) YING-YANG I;	IKANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2
Coumpound		YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA;	CHAIN: A, B;	YY 1: CHAIN: C: ADENO-	ASSOCIATED VIRUS PS	CHAIN: A, B;			YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS P5	INITIATOR ELEMENT DNA;	CHAIN: A, B;		,	YY1: CHAIN: C: ADENO-	ASSOCIATED VIRUS P5	INITIATOR ELEMENT DNA;	CHAIN: A, B;			YY1; CHAIN: C; ADENO-	ASSOCIATED VIRUS P5	CITABLE BEEMEN UNA;	CHAIN: A, B;			YY1; CHAIN: C; ADENO-	ASSOCIATED VIRUS PS	CHAIN: A, B;
SeqFold score																												
PMF score		_		-					0.88					-						. 66.0						_		
Verify score		0.23		0.27					0.04					0.2						-0.09						-0.06		
PSI- BLAST		1.80E-34		5.40E-34					6.50E-52					3.60E-34						7.20E-36						1.40E-34		
End		371		399					483					455						483						115		
Start		270	,	298	}				2	_				354						382		_				410		
Chain 19		U .		U	•				ပ					ပ						ပ						ပ		
PDB		Iubd		Inbd				1	Pappl					lubd						lubd				_		1ubd		
SEQ SO B		832		832					832					832						832						832		

PDB annotation	FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1;	IRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2	FINGER PROTEIN, DNA-PROTEIN	(TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION	TRANSCRIPTION INITIATION.	INITIATOR ELEMENT, YY1, ZINC 2	FINGER PROTEIN, DNA-PROTEIN	RECOGNITION, 3 COMPLEX	COMPLEY (TO ANCODEDITION	REGULATIONONA) YING-YANG 1:	TRANSCRIPTION INITIATION,	INITIATOR ELEMENT, YY1, ZINC 2	FINGER PROTEIN, DNA-PROTEIN	RECOGNITION, 3 COMPLEX	(TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA) YING-YANG 1;	INCLUSION INTERNATION,	FINGER PROTEIN. DNA-PROTEIN	RECOGNITION, 3 COMPLEX	(TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA) YING-YANG 1;	IKANSCRIPTION INITIATION,	TRICES DE CEEMENI, YYI, ZINC 2	PINGER PROTEIN, UNA-PROTEIN	TRANSCRIPTION REGILIATION/DNA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION,
Coumpound		YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS PS	INITIATOR ELEMENT DNA; CHAIN: A, B;			YY1; CHAIN: C; ADENO-	INITIATOR ELEMENT DNA:	CHAIN: A, B;			VVI. CHARI. C. ADENO	ASSOCIATED VIRUS P5	INITIATOR ELEMENT DNA;	CHAIN: A, B;				YY1; CHAIN: C; ADENO-	ASSOCIATED VIRUS PS	CHAIN A B.	(2,1,1,1,1,1,1,1,1,1,1,1,1,1,1,1,1,1,1,1			YY1; CHAIN: C; ADENO-	ASSOCIATED VIRUS PS	CITATION ELEMENT DNA;	CHAIN: A. B;			YY1; CHAIN: C; ADENO-	ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA;
SeqFold score																														86.51	
PMF	9	0.53				0.34	•									-								0.98							
Verify		-0.42				0.02					0.03	3						0.22						0.43							
PSI- BLAST		3.90E-42				3.60E-32					1 305-47	11000	-					5.40E-36						5.20E-51	_				:	5.20E-51	
End		594				999					623	3						622						829						619	
Start		428				466					\$10	}						522					ļ	267						569	
Chain ID		U				ပ					ļ	,						ບ						ပ ပ						၁	
PDB ID		pqn1				pqnl					74"[3						lubd		_				Inpq						Inbd	
SEQ No.		832		_		832					833	3					-	832						832	_		_			832	

PDB annotation	INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	ZINC FINGER DNA BINDING DOMAIN DNA BINDING MOTIF, ZINC FINGER DNA BINDING DOMAIN	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI, GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
Coumpound	CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI, CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	SWIS; CHAIN: NULL;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;
SeqFold score							-		
PMF		-	0.18	0.72	0.46	0.49	_		-
Verify score		90.0	6.04 4.04	0.15	-0.16	-0.24	-0.21	1.	0.06
PSI- BLAST	١	5.40E-34	2.60E-06	0.00013	1.30E-36	1.60E-31	2.60E-52	5.20E-63	3.90E-64
End		829	089	089	317	286	345	401	429
Start	_	577	159	653	133	150	208	235	262
Chain D		U	U		4	٧	٧	∢	Ą
PDB		Iubd	lubd	1zfd	2gli	2gli	2gli	2gli	2gli
SEQ B		832	832	832	832	832	832	832	832

					,		,				
PDB annotation	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	HYDROLASE HYDROLASE, HALOALKANE DEHALOGENASE, AI BHARETA LYCHOM ASE	HYDROLASE A/B HYDROLASE FOLD,
Conmpound	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZNC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA: CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	HALOALKANE DEHALOGENASE; CHAIN: NITT:	HALOALKANE
SeqFold score									91.33		
PMF	-	0.99	0.19	0.99	0.49	0.19	0.99			0.12	0.21
Verify score	0.01	0.25	-0.4	0.03	-0.21	-0.07	-0.02	0.53		-0.1	-0.19
PSI- BLAST	3.60E-33	2.60E-64	3.90E-57	3.60E-34	1.10E-32	5.20E-60	5.40E-31	3.90E-64	3.90E-64	0.00078	0.0013
End AA	398	485	965	482	265	652	624	678	089	1332	1332
Start AA	270	318	346	354	438	459	494	541	541	1240	1240
Chain D	∢	<	∢	∢	∢	<	<	4	4		Ą
PDB CI	2gli	2gli	2gli	2gli	2gli	2gli	2gli	2gli	2gli ·	1b6g	Icqw
SEQ B B S	832	832	832	832	832	832	832	832	832	834	834

		ol- Ion,		OL NAS,	 ניז	T	S		SE, 1	DENT		ASE,	ä		ER	,
ation	GNO	LGLYCER(2 E, OXYAN ASE	SETA PROTEIN	L-GLYCER IA-BETA SEUDOMO BITOR	ATHESIS S-LYASE; ATHESIS, S-LYASE; PHATE, N BINDIN	104 0776	REDOXAL	OLD	RANSFER	LP-DEPEN NE	SERINE PI P	ransfer	EPENDENT	A LYASE	EPENDENT	
PDB annotation	AASE I-S B	; TRIACYI ; X-RAY GRAPHY, VADACEA , HYDROI	ALPHA/E	TRIACYI ASE, ALPH FOLD, PS TE 2 INHI	N BIOSYN N BIOSYN N BIOSYN N INDOL! 2 5'-PHOS	OT TO	ROLE, PY	IA/BETA I	SE SHMT; ETHYL TI TABOLIS	LYASE, L JETHIONI SIS	SE SHMT,	AMINO 1	SE PLP-DI	C-S 2 BET	SE PLP-DI	
	DEHALOGENASE I-S BOND	HYDROLASE TRIACYLGLYCEROL- HYDROLASE, X-RAY CRYSTALLOGRAPHY, 2 PSEUDOMONADACEAE, OXYANION, CIS-PEPTIDE, HYDROLASE	HYDROLASE ALPHABETA HYDROLASE, GLYCOPROTEIN	HYDROLASE TRIACYL-GLYCEROL LIPASE, LIPASE, ALPHA-BETA HYDROLASE FOLD, PSEUDOMONAS, PHOSPHONATE 2 INHIBITOR	TRYPTOPHAN BIOSYNTHESIS TRYPTOPHAN INDOLE-LYASE; TRYPTOPHAN BIOSYNTHESIS, TRYPTOPHAN INDOLE-LYASE, PYRIDOXAL 2 5-PHOSPHATE, MONOVALENT CATION BINDING	THE ANIGHER A ST. TR ANIGHTR ASE	METABOLIC ROLE, PYRIDOXAL 5'- PHOSPHATE	LYASE ALPHA/BETA FOLD	TRANSFERASE SHMT; HYDROXYMETHYL TRANSFERASE, 1 CARBON METABOLISM	LYASE CGS; LYASE, LLP-DEPENDENT ENZYMES, METHIONINE BIOSYNTHESIS	TRANSFERASE SHMT, SERINE	ASPARTATE, AMINO TRANSFERASE, (AATHIKE FOLD)	TRANSFERASE PLP-DEPENDENT FN7YMES IRON-SHI FIR-CHISTER	SYNTHESIS, C-S 2 BETA LYASE	TRANSFERASE PLP-DEPENDENT ENZYMES, IRON-SULFUR-CLUSTER	
	AIN: A;	: NULL;	IAIN: A;	<u>ដ</u> ្ឋ	HAIN:	Ī	RANSF	IN: A:	RANSF	MMA- 4, B, C,	DANCE	C, D	E,		ű	
Coumpound	NASE; 1-	LYCEROL IE; CHAIN	L PROTEIN ASE 1; CF	NG LIPAS	ANASE; C		VETHYLT AIN: A:	EIN; CHA	METHYLT AIN: A. B.	CHAIN:	TIMAL	AIN: A, B,	NSFERAS	•	NSFERAS	
රී	DEHALOGENASE; 1- CHLOROHEXANE CHAIN: A;	TRIACYLGLYCEROL HYDROLASE; CHAIN: NULL;	PALMITOYL PROTEIN THIOESTERASE 1; CHAIN: A;	LACTONIZING LIPASE; CHAIN: A;	TRYPTOPHANASE; CHAIN: A, B, C, D;	THE PLANT	SEKURE HYDROXYMETHYLTRANSF ERASE: CHAIN: A:	CSDB PROTEIN; CHAIN: A;	SERINE HYDROXYMETHYLTRANSF ERASE: CHAIN: A. B:	CYSTATHIONINE GAMMA- SYNTHASE; CHAIN: A, B, C, D:	SERINE HYDROXYMETHYI TBANSE	ERASE; CHAIN: A, B, C, D:	AMINOTRANSFERASE;	Circuit: 0.	AMINOTRANSFERASE; CHAIN: A. B:	
SeqFold score				·												_
PMF		0.69	0.53	0.46	90.0	200	,0.0 -	-1.41	-0.13	0	-1.41		-		-	_
Verify score		0.04	-0.1	0.13	0.23	000	0.00	80.0	0.05	-0.28	0.17		0.22		0.1	_
PSI- BLAST		0.00026	0.00026	0.00026	5.40E-07	2000	05-207-7	9.00E-41	7.20E-37	1.60E-31	3.60E-40		1.30E-50		1.80E-45	
End		1362	1362	1362	241	,;;	047	244	246	242	252		248		248	
Start		1251	1251	1251	<u>E</u>	:	2	7	12	4	4		_		-	
Chain ID			⋖	¥.	₹	-	€	V	∢	4	٧		V		¥	
PDB DD		lcvl	lei9	lex9	lax4		104	1c0n	1cj0	lcs1	1dfo		leg5		legS	
SEQ EQ		834	834	834	836	926	8	836	836	836	836		836		836	_

	Т	, 	1		 7			ı		
PDB annotation	TRANSFERASE SHMT; SERINE- GLYCINE CONVERSION, PYRIDOXAL 5-PHOSPHATE, 2 TETRAHYDROFOLATE, ASYMMETRIC DIMER	LYASE FES CLUSTER BIOSYNTHESIS, PYRIDOXAL 5'-PHOSPHATE, 2 THIOCYSTEINE, AMMOACRYLATE, ENZYME-PRODUCT COMPLEX	CARBOXYLIC ESTER HYDROLASE PHOSPHOLIPASE, TRIMER, CALCIUM BINDING, ACTIVATOR SITE, 2 CARBOXYLIC ESTER HYDROLASE	HYDROLASE PLA2, PHOSPHATIDE SN-2 ACYLHYDROLASE; HYDROLASE; PHOSPHOLIPASE A2, LIPID DEGRADATION, PRESYNAPTIC 2 NEUROTOXIN, VENOM	HYDROLASE HYDROLASE, PHOSPHOLIPASE A2, PLATELET AGGREGATION INHIBITOR, 2 PBPB HEADER MODRES					LIPID DEGRADATION PHOSPHOLIPASE A2, LIPID
Coumpound	SERINE HYDROXYMETHYLTRANSF ERASE; CHAIN: A, B, C, D;	L-CYSTEINEL-CYSTINE C-S LYASE; CHAIN: A, B;	PHOSPHOLIPASE A2; CHAIN: NULL;	PHOSPHOLIPASE A2; CHAIN: NULL;	PHOSPHOLIPASE A2; CHAIN: NULL;	HYDROLASE PHOSPHOLIPASE A2 (E.C.3.1.14) 1POA 3	HYDROLASE PHOSPHOLIPASE A2 (E.C.3.1.1.4) COMPLEX WITH THE 1POC 3 TRANSITION- STATE ANALOGUE 1POC 4	HYDROLASE PHOSPHOLIPASE A2 E.C.3.1.1.4) COMPLEX WITH THE 1POC 3 TRANSITION- STATE ANALOGUE 1POC 4	HYDROLASE CALCIUM- FREE PHOSPHOLIPASE A=2= (E.C.3.1.1.4) 1PP2 4	PHOSPHOLIPASE A2; CHAIN: A, B;
SeqFold score							99.59			
PMF score	-0.11	0.83	-0.08	-0.18	-0.13	-0.14		0.21	-0.14	-0.17
Verify score	0.41	-0.03	0.38	0.07	0.07	0.3		0.08	0.2	0.12
PSI- BLAST	7.20E-35	3.90E-28	4.80E-39	4.80E-41	1.60E-40	1.60E-38	1.40E-36	0.0054	1.60E-38	3.20E-44
End	246	241	234	244	253	234	289	438	253	253
Start	12	-	131	131	131	131	151	355	. 131	131
Chain ID	4	4							~	Ą
PDB	leji	lelu	1a3d	1ae7	15k9	Ipoa	Ipoc	lpoc	1pp2	lvap
SEQ NO.	836	836	837	837	837	837	837	837	837	837

PDB annotation	Н		HYDROLASE HYDROLASE, LIPID DEGRADATION, CALCIUM, PRESYNAPTIC 2 NEUROTOXIN, VENOM		OXIDOREDUCTASE PDZ DOMAIN, NNOS, NITRIC OXIDE SYNTHASE	PEPTIDE RECOGNITION PEPTIDE RECOGNITION, PROTEIN LOCALIZATION	PEPTIDE RECOGNITION PEPTIDE RECOGNITION, PROTEIN LOCALIZATION	CYTOKINE LCF, CYTOKINE, LYMPHOCYTE CHEMOATTRACTANT FACTOR, PDZ DOMAIN	CYTOKINE LCF, CYTOKINE, LYMPHOCYTE CHEMOATTRACTANT FACTOR, PDZ DOMAIN	KINASE HCASK, GLGF REPEAT, DHR; PDZ DOMAIN, NEUREXIN, SYNDECAN, RECEPTOR CLUSTERING,	SIGNAL TRANSDUCTION HDLG, DHR3 DOMAIN; SIGNAL
Coumpound		PHOSPHOLIPASE A2; CHAIN: NULL;	PHOSPHOLIPASE A2; CHAIN: A, B;	HYDROLASE(CARBOXYL ESTER) PHOSPHOLIPASE A=2=(PHOSPHATIDE-2- ACYL-HYDROLASE) MUTANT 3P2P 4 WITH ASP 59 REPLACED BY SER, SER 60 REPLACED BY GY, 3P2P REPLACED BY TYR 3P2P 6 (D59S\$, (S60G\$, DEL(62-66), N67Y\$) (E.C.3.1.4) 3P2P 7	NEURONAL NITRIC OXIDE SYNTHASE; CHAIN: A; HEPTAPEPTIDE; CHAIN: B;	PSD-95; CHAIN: A; CRIPT; CHAIN: B;	PSD-95; CHAIN: A; CRIPT; CHAIN: B;	NTERLEUKIN 16; CHAIN: NULL;	NULL;	HCASK/LIN-2 PROTEIN; CHAIN: A, B;	HUMAN DISCS LARGE PROTEIN: CHAIN: NULL:
SeqFold score	1										
PMF score		-0.11	-0.08	-0.06	86.0	0.72	0.1	0.99	0.45	99.0	0.88
Verify score		0.15	0.4	0.34	0.79	0.21	-0.25	0.59	-0.03	0.04	0.45
PSI- BLAST		4.80E-41	1.30E-40	1.60E-38	1.30E-13	1.10E-18	6.40E-10	7.20E-16	1.40E-14	1.30E-14	6.40E-17
End AA		248	244	236	106	102	317	109	108	108	113
Start		131	131	132	21	12	254	21	4	21	20
Chain ID			4	∢	4	¥	4			4	
PDB		lvip	2not	3р2р	1 b8q	1be9	1be9	1116	1116	lkwa	1 pdr
SEQ EQ		837	837	837	838	838	838	838	838	838	838

						_,							_	
PDB annotation	TRANSDUCTION, SH3 DOMAIN, REPEAT	OXIDOREDUCTASE BETA-FINGER	MEMBRANE PROTEIN/OXIDOREDUCTASE BETA- FINGER, HETERODIMER	PEPTIDE RECOGNITION PSD-95; PDZ DOMAIN, NEURONAL NITRIC OXIDE SYNTHASE, NMDA RECEPTOR 2 BINDING	HYDROLASE PDZ DOMAIN, HUMAN PHOSPHATASE, HPTP1E, PTP-BAS, SPECIFICITY 2 OF BINDING		OXIDOREDUCTASE (D, L) STEREOSPECIFIC OPINE	DEHYDROGENASE, OXIDOREDUCTASE	OXIDOREDUCTASE OXYDOREDUCTASE	TRANSFERASE TRANSFORMYLASE, PURINE BIOSYNTHESIS, ATP-GRASP	OXIDOREDUCTASE LYSINE BIOSYNTHESIS, ALPHA- AMINOADIPATE PATHWAY, 2 SACCHAROPINE REDUCTASE, DEHYDROGENASE	OXIDOREDUCTASE OXIDOREDUCTASE, OXIDOREDUCTASE, NAD		TRANSFORMING GROWTH FACTOR OSTEOGENIC PROTEIN-1, HOP-1,
Coumpound		NEURONAL NITRIC OXIDE SYNTHASE (RESIDUES 1- 130); CHAIN: A;	ALPHA-1 SYNTROPHIN (RESIDUES 77-171); CHAIN: A; NEURONAL NITRIC OXIDE SYNTHASE (RESIDUES 1-130); CHAIN: B;	POSTSYNAPTIC DENSITY PROTEIN 95; CHAIN: A;	TYROSINE PHOSPHATASE (PTP-BAS, TYPE 1); CHAIN: A;		N-(I-D-CARBOXYLETHYL)- L-NORVALINE	DEHYDROGENASE; CHAIN:	GLYCERALDEHYDE-3- PHOSPHATE DEHYDROGENASE, CHAIN: P. R. O. O.	PHOSPHORIBOSYLGLYCINA MIDE FORMYLTRANSFERASE 2; CHAIN: A. B:	SACCHAROPINE REDUCTASE; CHAIN: A;	L-ALANINE DEHYDROGENASE; CHAIN: A;		BONE MORPHOGENETIC PROTEIN-7: CHAIN: NULL:
SeqFold score										•				
PMF		0.57	0.86 –		0.45		0.17		0.09	0.05	-	0.49		_
Verify score		-0.12	0.49	0.28	-0.09		-0.18	·	-0.1	-0.09	1.04	-0.24		0.14
PSI- BLAST		9.00E-15	3.20E-17	1.40E-18	6.40E-17		1.60E-05		0.0069	0.0054	4.80E-58	1,30E-61		9.60E-48
End		106	103	107	66		594		570	599	924	460		213
Start		21	20	20	23		481		480	482	481	25		112
Chain		4	∢	₹	A				<u>a</u>	4	∢	4		
PDB DD		Iqau	Iqav	1qlc	3pdz		1bg6		1c2	leyz	169	1pjc		1bm n
SEQ	Ö	838	838	838	838		840		840	840	840	840		842

PDB annotation	PTSI PROTEIN-PEPTIDE COMPLEX, TETRATRICOPEPTIDE REPEAT, TPR, 2 HELICAL REPEAT	SIGNALING PROTEIN PEROXISMORE RECEPTOR 1, PTSI-BP, PEROXIN-5, PTSI PROTEIN-PEPTIDE COMPLEX, TETRATRICOPEPTIDE REPEAT, TPR, 2 HELICAL REPEAT	SIGNALING PROTEIN PEROXISMOKE RECEPTOR 1, PTSI-BP, PEROXIN-5, PTSI PROTEIN-PEPTIDE COMPLEX, TETRATRICOPEPTIDE REPEAT, TPR, 2 HELICAL REPEAT	SIGNALING PROTEIN PEROXISMORE RECEPTOR 1, PTS1-BP, PEROXIN-5, PTS1 PROTEIN-PEPTIDE COMPLEX, TETRATRICOPEPTIDE REPEAT, TPR, 2 HELICAL REPEAT	TRANSFERASE PROTO-ONCOGENE TYROSINE KINASE; PROTO- ONCOGENE, TRANSFERASE, TYROSINE-PROTEIN (MASE, 2 PHOSPHORYLATION, ATP-BINDING, MYRISTYLATION, SH3 DOMAIN, 3 COMPLEX (PHOSPHOTRANSFERASE/PEPTIDE)		COMPLEX (KINASE/PEPTIDE)	- COMPLEX (TRANSFERASE/PEPTIDE)
Coumpound	CHAIN: A, B; PTS1- CONTAINING PEPTIDE; CHAIN: C, D;	PEROXISOMAL TARGETING SIGNAL I RECEPTOR; CHAIN: A, B; PTS1- CONTAINING PEPTIDE; CHAIN: C, D;	PEROXISOMAL TARGETING SIGNAL 1 RECEPTOR; CHAIN: A, B; PTS1-CONTAINING PEPTIDE; CHAIN: C, D;	PEROXISOMAL TARGETING SIGNAL I RECEPTOR; CHAIN: A, B; PTS1- CONTAINING PEPTIDE; CHAIN: C, D;	PHOSPHOTRANSFERASE FYN; CHAIN: A; 3BP-2; CHAIN: B;	SIGNAL TRANSDUCTION PROTEIN GROWTH FACTOR RECEPTOR-BOUND PROTEIN 2 (GRB2, N-TERMINAL IGBR 3 SH3 DOMAIN) COMPLEXED WITH SOS-A PEPTIDE IGBR 4 (NMR, 29 STRUCTURES) IGBR 5	P56—LCK— TYROSINE KINASE, ILCK 7 CHAIN: A; ILCK 8 TAIL PHOSPHOPETIDE TEGQ(PHOSPHO)YOPOPA; ILCK 14 CHAIN: B; ILCK 15	C-SRC: CHAIN: C; NL1 (MN7-
SeqFold score								
PMF score		0.62	-0.15	0.99	6.4	0.59	0.28	0 29
Verify score		0.29	90.0	0.02	0.23	0.74	0.09	40
PSI- BLAST		3.60E-09	1.30E-20	1.10E-12	1.80E-07	1.40E-06	1.80E-05	1 ROF-07
End		804	669	842	242	249	788	242
Start		445	452	576	193	181	193	103
Chain		4	¥	V	∢	. ✓	∢	ر
PDB ID		1fch	1fch	1fgh	1fyn	1gbr	11ck	120
SEQ	Ö	843	843	843	843	843	843	843

PDB annotation	SRC, SH3 DOMAIN, LIGANDS, NON- PEPTIDE ELEMENTS, 2 COMPLEX (TRANSFERASE/PEPTIDE)			TYROSINE KINASE TYROSINE KINASE-INHIBITOR COMPLEX, DOWN-REGULATED KINASE, 2 ORDERED ACTIVATION LOOP	PROTEIN TRANSPORT HELIX-TURN- HELIX TPR-LIKE REPEAT, PROTEIN TRANSPORT	PROTEIN TRANSPORT HELIX-TURN- HELIX TPR-LIKE REPEAT, PROTEIN TRANSPORT		TRANSFERASE HCK; SH3, PROTEIN TYROSINE KINASE, SIGNAL TRANSDUCTION, 2 TRANSFERASE	IMMUNOGLOBULIN IMMUNOGLOBULIN	IMMUNOGLOBULIN IMMUNOGLOBULIN, FAB FRAGMENT, HUMANISATION	COMPLEX (VIRAL CAPSID/IMM/UNOGLOBULIN) HIV-1 CA, HIV CA, HIV P24, P24; FAB, FAB LIGHT CHAIN, FAB HEAVY CHAIN COMPLEX (VIRAL CAPSID/IMM/UNOGLOBULIN), HIV,
Coumpound	MNZ-MNI-PLPPLP); CHAIN: N;	PHOSPHOTRANSFERASE PHOSPHATIDYLINOSITOL 3- KINASE (P85-ALPHA STIBILITY DENT 3 SH3	DOMAIN) (NMR, MINIMIZED AVERAGE STRUCTURE) IPNI 4	HAEMATOPOETIC CELL KINASE (HCK); CHAIN: A;	VESICULAR TRANSPORT PROTEIN SEC17; CHAIN: A;	VESICULAR TRANSPORT PROTEIN SEC17; CHAIN: A;	PHÖSPHOTRANSFERASE FYN PROTO-ONCOGENE TYROSINE KINASE (E.C.2.7.1.112) ISHF 3 (SH3 DOMAIN) ISHF 4	HEMATOPOIETIC CELL KINASE; CHAIN: NULL;	2E8 (IGG1=KAPPA=) ANTIBODY; CHAIN: L, H, M, P:	ANTIBODY CTM01; CHAIN: L, H;	HUMAN IMMUNODEFICIENCY VIRUS TYPE I CAPSID CHAIN: A. B; ANTIBODY FAB25.3 FRAGMENT; CHAIN: H. K. L. M:
SeqFold score											
PMF		0.17		0.17	0.15	. 60.0	0.54	0.89	-0.06	0.03	0.24
Verify score		0.54	-	0.09	90.0	-0.03	0.38	0.45	0.08	-0.18	0.05
PSI- BLAST		9.00E-10		3.60E-07	1.80E-07	8.00E-06	9.00E-08	9.00E-06	6.40E-51	4.80E-53	4.80E-63
End AA		249		282	573	1101	242	255	414	599	220
Start AA		182		193	418	949		193	236	419	21
Chain				∢	V	¥	∢		н	н	H
PDB		Ipnj		lqcf	lqqe	lgge	1shf	4hck	1.20 E+09	1ae6	lafv
SEQ B B		843		843	843	843	843	843	846	846	846

PDB annotation	CAPSID PROTEIN, 2 P24	INSECT IMMUNITY INSECT IMMUNITY, LPS-BINDING, HOMOPHILIC ADHESION	INSECT IMMUNITY INSECT IMMUNITY, LPS-BINDING, HOMOPHILIC ADHESION	INSECT IMMUNITY INSECT IMMUNITY, LPS-BINDING, HOMOPHILIC ADHESION	INSECT IMMUNITY INSECTIMMUNITY, LPS-BINDING, HOMOPHILIC ADHESION	INSECT IMMUNITY INSECTIMMUNITY, LPS-BINDING, HOMOPHILIC ADHESION	RECEPTOR RECEPTOR, SIGNAL TRANSDUCER OF IL-6 TYPE TYPOKINES, THIRD 2 IN-TERMINAL DOMAIN, TRANSMEMBRANE, GLYCOPROTEIN	IMMUNE SYSTEM IMMUNOGLOBULIN, IMMUNE SYSTEM	IMMUNOGLOBULIN IMMUNOGLOBULIN, FAB COMPLEX, IDIOTOPE, ANTI-IDIOTOPE	IMMUNE SYSTEM IMMUNOGLOBULIN, IGG1; IMMUNOGLOBULIN, IGG1; IMMUNOGLOBULIN, IGG1; RAGMENT, CROSS-REACTIVITY, HIVI PROTEASE, ENZYME 2 INHIBITION, IMMUNOGLOBULIN	IMMUNE SYSTEM ANTI-PRION FAB
Соитроинд		HEMOLIN; CHAIN: A, B;	HEMOLIN: CHAIN: A. B;	HEMOLIN; CHAIN: A, B;	HEMOLIN; CHAIN: A, B;	HEMOLIN; CHAIN: A, B;	GP130; CHAIN: NULL;	MONOCLONAL ANTIBODY MRK-16 (LIGHT CHAIN); CHAIN: A, C; MONOCLONAL ANTIBODY MRK-16 (HEAVY CHAIN); CHAIN: B, D;	IG HEAVY CHAIN V REGIONS; CHAIN: A; IG HEAVY CHAIN: A; IG CHAIN: B; IG HEAVY CHAIN V REGIONS; CHAIN: C; IG HEAVY CHAIN: C; IG CHAIN: D;	IGGI ANTIBODY 1696 (LIGHT CHAIN); CHAIN: L; IGGI ANTIBODY 1696 (VARIABLE HEAVY CHAIN); CHAIN: H; IGGI ANTIBODY 1696 (CONSTANT HEAVY CHAIN); CHAIN: I;	FAB ANTIBODY LIGHT
SeqFold score		159.92				·			,		
PMF			_	0.54	0.66	0.21	-0.05	0.04	-0.13	-0.19	-0.11
Verify score			0.28	0.29	0.15	0.14	0.07	0.36	0.17	0.03	90.0
PSI- BLAST		5.40E-62	5.40E-62	1.60E-31	7.20E-48	3.20E-22	1.40E-12	3.20E-64	1.60E-49	6.40E-23	3.20E-49
End		503	503	493	404	713	597	220	412	603	415
Start		134	135	138	27	323	208	21	235	223	236
Chain		A	4	4	¥	4		m	α	н	H
PDB EDB	1	1bih	1bih	1bih	1bih	1bih	1bj8	1bln	1cic	1c17	1cr9
SEQ ID	ÿ	846	846	846	846	846	846	846	846	846	846

																	_					_					
PDB annotation	3F4; ANTI-PRION FAB 3F4 ANTI-PRION ANTIBODY, FAB 3F4	CELL ADHESION NEURAL CELL ADHESION	CELL ADHESION NEURAL CELL ADHESION	CELL ADHESION NEURAL CELL ADHESION	CELL ADHESION NEURAL CELL ADHESION	CELL ADHESION NEURAL CELL ADHESION	CELL ADHESION NEURAL CELL ADHESION	CELL ADHESION NEURAL CELL ADHESION	CELL ADHESION NEURAL CELL ADHESION	GROWTH FACTOR/GROWTH FACTOR	RECEPTOR FGF, FGFR,	IMMUNOGLOBULIN-LIKE, SIGNAL	TRANSDUCTION, 2 DIMERIZATION,	GROWTH FACTOR/GROWTH FACTOR RECEPTOR	GROWTH FACTOR/GROWTH FACTOR	RECEPTOR FGF, FGFR,	IMMUNOGLOBULIN-LIKE, SIGNAL	GROWTH FACTOR/GROWTH FACTOR	RECEPTOR	GROWTH FACTOR/GROWTH FACTOR	RECEPTOR FGF, FGFR,	TRANSOLICATION 2 DIMERIZATION	GROWTH FACTOR/GROWTH FACTOR	RECEPTOR	GROWTH FACTOR/GROWTH FACTOR	RECEPTOR FGF, FGFR,	TRANSDUCTION, 2 DIMERIZATION,
Coumpound	CHAIN; CHAIN: L; FAB ANTIBODY HEAVY CHAIN; CHAIN: H;	AXONIN-1; CHAIN: A;	AXONIN-1; CHAIN: A;	AXONIN-1; CHAIN: A;	AXONIN-1; CHAIN: A;	AXONIN-1; CHAIN: A;	AXONIN-1; CHAIN: A;	AXONIN-1; CHAIN: A;	AXONIN-1; CHAIN: A;	FIBROBLAST GROWTH	FACTOR 2; CHAIN: A, B;	FIBROBLAST GROWTH	FACTOR RECEPTOR 1;	CHAIN: C, D;	FIBROBLAST GROWTH	FACTOR 2; CHAIN: A, B;	FIBROBLAST GROWTH	CHAIN: C, D;		FIBROBLAST GROWTH	FACTOR 2; CHAIN: A, B;	FIBRUBLASI GROWIN	CHAIN: C. D.		FIBROBLAST GROWTH	FACTOR 2; CHAIN: A, B;	FACTOR RECEPTOR 1;
SeqFold score																								-			
PMF score		-	-	0.43	-0.14	96.0	1	0.58	0.51	0.94					0.22					0.99					-0.03		
Verify score		80.0	0.24	-0.14	0.02	0.23	0.13	0.07	0.07	0.15					0.07					0.22					0.26		
PSI- BLAST		3.60E-59	4.80E-45	1.60E-47	1.10E-37	1.80E-44	6.40E-42	1.60E-53	3.20E-33	1.80E-30					3.20E-21					1.80E-32					8.00E-21		
End AA		503	503	321	412	592	602	404	713	320					601					320					109		
Start AA	•	135 ·	138	2	20	226	236	24	320	135					430					135					430		
Chain The Chain		٧	A	¥	V	4	٧	V	¥	U					U							_			α		
EDB CI		lcs6	1036	1036	1cs6	1cs6	1cs6	1cs6	lcs6	lcvs					Icvs					lcvs	•				lcvs		
SEQ RO:		846	846	846	846	846	846	846	846	846					846					846					846		

PDB annotation	GROWTH FACTOR/GROWTH FACTOR RECEPTOR	VIRUS/VIRAL, PROTEIN, RECEPTOR CD155, PVR, HUMAN POLIOVIRUS, ELECTRON MICROSCOPY, 2 POLIOVIRUS-RECEPTOR COMPLEX, VIRUS/VIRAL PROTEIN, RECEPTOR	VIRUS/VIRAL PROTEIN, RECEPTOR CD155, PVR, HUMAN POLIOVIRUS, ELECTRON MICROSCOPY, 2 POLIOVIRUS-RECEPTOR COMPLEX, VIRUS/VIRAL PROTEIN, RECEPTOR	VIRUS/VIRAL PROTEIN, RECEPTOR CD155, PVR, HUMAN POLIOVIRUS, EELECTRON MICROSCOPY, 2 POLIOVIRUS-RECEPTOR COMPLEX, VIRUS/VIRAL PROTEIN, RECEPTOR	IMMUNE SYSTEM FC IGG PHAGE DISPLAY PEPTIDE	IMMUNE SYSTEM IMMUNOGLOBULIN FOLD, ANTIBODY, IGM, FV	COMPLEX (ANTIBODY ANTIGEN) 1,4- BETA-N-ACETYLMURAMIDASE C; SINGLE-DOMAIN ANTIBODY, TURKEY EGG-WHITE LYSOZYME, 2 ANTIBODY-PROTEIN COMPLEX, SINGLE-CHAIN FV FRAGMENT	CELL ADHESION NCAM; NCAM, IMMUNOGLOBULIN FOLD, GLYCOPROTEIN	CELL ADHESION NCAM; NCAM, IMMUNOGLOBULIN FOLD, GLYCOPROTEIN	GROWTH FACTOR/GROWTH FACTOR RECEPTOR FGF2; FGFR2; IMMUNOGLOBULIN (IG)LIKE
Coumpound	CHAIN: C, D;	POLIOVIRUS RECEPTOR; CHAIN: R; VP1; CHAIN: 1; VP2; CHAIN: 2; VP3; CHAIN: 3; VP4; CHAIN: 4;	Ë	POLIOVIRUS RECEPTOR; CHAIN: R; VPI; CHAIN: I; VP2; CHAIN: 2; VP3; CHAIN: 3; VP4; CHAIN: 4;	IMMUNOGLOBULIN LAMBDA HEAVY CHAIN; CHAIN: A, B; ENGINEERED PEPTIDE; CHAIN: E, F;	HAIN:	SCFV FRAGMENT 1F9; CHAIN: A, B; TURKEY EGG- WHITE LYSOZYME C; CHAIN: X, Y;	NEURAL CELL ADHESION MOLECULE; CHAIN: A, B, C, D;	NEURAL CELL ADHESION MOLECULE; CHAIN: A, B, C, D;	FIBROBLAST GROWTH FACTOR 2; CHAIN: A, B, C, D; FIBROBLAST GROWTH
SeqFold score										
PMF		0.65	0.21	0.01	0.05	-0.09	-0.03	6.0	0.78	96.0
Verify score		-0.3	-0.31	-0.4	-0.27	0.09	0.14	0.34	0.37	0.07
PSI- BLAST		7.20E-36	1.10E-38	3.60E-44	8.00E-40	3.20E-38	6.40E-42	1.80E-31	1.30E-24	1.40E-34
End		404	503	319	411	130	206	319	503	320
Start AA		141	228	78	234	21	21	142	324	135
Chain		A.	&	æ	V	н	∢	∀	∀	ப
PDB U		Idgi	1dgi	1dgi	1dn2	1dql	1dzb	lepf	lepf	lev2
SEQ.	2	846	846	846	846	846	846	846	846	846

PDB annotation	DOMAINS BELONGING TO THE I-SET 2 SUBGROUP WITHIN IG-LIKE DOMAINS, B-TREFOIL FOLD	GROWTH FACTOR/GROWTH FACTOR	MACEFIOR FORZ; FORZ;	DOMAINS BELONGING TO THE I-SET	2 SUBGROUP WITHIN IG-LIKE	DOMAINS, B-TREFOIL FOLD	GROWTH FACTOR/GROWTH FACTOR	RECEPTOR FGF2; FGFR2;	DOMATING BELONGING TO THE L-SET	2 SUBGROUP WITHIN IG-LIKE	DOMAINS, B-TREFOIL FOLD	GROWTH FACTOR/GROWTH FACTOR	RECEPTOR FGF2; FGFR2;	IMMUNOGLOBULIN (IG)LINE	DOMAINS BELONGING TO THE 1-SET	DOMAINS B-TREFOIL FOLD	GPOWTH FACTOR/GPOWTH FACTOR	RECEPTOR FGF2: FGFR2:	IMMUNOGLOBULIN (IG)LIKE	DOMAINS BELONGING TO THE I-SET	2 SUBGROUP WITHIN IG-LIKE	DOMAINS, B-TREFOIL FOLD	GROWTH FACTOR/GROWTH FACTOR	RECEPTOR FGF1; FGFR1;	IMMUNOGLOBULIN (IG) LIKE	DOMAINS BELONGING 10 THE 1-SET	2 SUBGROUP WITHIN IG-LIKE	GROWTH FACTOR/GROWTH FACTOR	RECEPTOR FGF1; FGFR1;	IMMUNOGLOBULIN (IG) LIKE	DOMAINS BELONGING TO THE I-SET	2 SUBGROUP WITHIN IG-LIKE	IMMUNE SYSTEM FC-EPSILON RI-	ALPHA; IMMUNOGLOBULIN FOLD,
Coumpound	FACTOR RECEPTOR 2; CHAIN: E, F, G, H;	FIBROBLAST GROWTH	FACIOR 2; CHAIN: A, B, C, D; FIRRORI AST GROWTH	FACTOR RECEPTOR 2:	CHAIN: E, F, G, H;		FIBROBLAST GROWTH	FACTOR 2; CHAIN: A, B, C, D;	FIBROBLASI GROWIN	CHAIN: E. E. G. H:		FIBROBLAST GROWTH	FACTOR 2; CHAIN: A, B, C, D;	FIBROBLASI GROWIN	FACTOR RECEPTOR 2;	CHAIN: E, F, G, H;	ETBDOBI AST CDOWTH	FACTOR 2: CHAIN: A B C D:	FIBROBLAST GROWTH	FACTOR RECEPTOR 2;	CHAIN: E, F, G, H;		FIBROBLAST GROWTH	FACTOR 1; CHAIN: A, B;	FIBROBLAST GROWTH	FACTOR RECEPTOR 1;	CHAIN: C, D;	FIBROBLAST GROWTH	FACTOR 1; CHAIN: A, B;	FIBROBLAST GROWTH	FACTOR RECEPTOR 1;	CHAIN: C, D;	HIGH AFFINITY	IMMUNOGLOBULIN
SeqFold score																																		
PMF		0.17					0.94		•			0.94					200	2					69.0					0.07					0.29	
Verify score		0.15					0.23					0.27					31.0	3					0.11					-0.07					0.09	
PSI- BLAST		7.20E-23					5.40E-33					3.60E-24					1 405-10	71-700:1					1.30E-30					6.40E-20					7.20E-28	
End		224					322					505					109	3					320					601					322	
Start AA		33					135					329					430	}					135					430					136	
Chain		Ξ					9					g					c	,					ပ					ပ					4	
PDB ID		lev2					lev2					lev2					10,0	1					1थ					levt					129	
SEQ NO		846					846					846			_	_	846	}					846					846					846	

PDB annotation	GLYCOPROTEIN, RECEPTOR, IGE- BINDING 2 PROTEIN	IMMUNE SYSTEM ANTI- CARBOHYDRATE ANTIBODY	IMMUNE SYSTEM HIGH AFFINITY IGE-FC RECEPTOR, FC(EPSILON) IGE- FC, IMMUNOGLOBULIN FOLD, GLYCOPROTEIN, RECEPTOR, IGE- BINDING 2 PROTEIN, IGE ANTIBODY, IGE-FC	HORMONEGROWTH FACTOR/HORMONE RECEPTOR 4- HELICAL BUNDLE, ALPHA HELICAL BUNDLE, TERNARY COMPLEX, FN 2 III DOMAINS, BETA SHEET DOMAINS, CYTOKINE-RECEPTOR COMPLEX		IMMUNE SYSTEM, MEMBRANE PROTEIN CD32; FC RECEPTOR, IMMUNOGLOULIN, LEUKOCYTE, CD32	HEPARIN AND INTEGRIN BINDING HEPARIN AND INTEGRIN BINDING	IMMUNE SYSTEM RECEPTOR BETA SANDWICH, IMMUNOGLOBULIN- LIKE, RECEPTOR			IMMUNOGLOBULIN INTACT IMMUNOGLOBULIN V REGION C
Coumpound	EPSILON RECEPTOR CHAIN: A;	ANTIBODY S-20-4, FAB FRAGMENT, LIGHT CHAIN; CHAIN: L; ANTIBODY S-20-4, FAB FRAGMENT, HEAVY CHAIN: CHAIN: H	HIGH AFFINITY IMMUNGGLOBULIN EPSILON RECEPTOR CHAIN: A; IG EPSILON CHAIN C REGION; CHAIN: B, D;	PLACENTAL LACTOGEN; CHAIN: A; PROLACTIN RECEPTOR; CHAIN: B, C;	IMMUNOGLOBULIN IMMUNOGLOBULIN FC AND FRAGMENT B OF PROTEIN A COMPLEX 1FC2 4	FC (GAMMA)RIIA; CHAIN: A;	FIBRONECTIN; CHAIN: A;	LOW AFFINITY IMMUNOGLOBULIN GAMMA FC REGION CHAIN: A;	T LYMPHOCYTE ADHESION GLYCOPROTEIN CD2 (HUMAN) IHNF 3	TLYMPHOCYTE ADHESION GLYCOPROTEIN CD2 (RAT) IHNG 3	IGG2A INTACT ANTIBODY - MAB231; CHAIN: A, B, C, D
SeqFold score											
PMF		0	0.18	0.03	0.07	0.23	0.52	0.36	90.0	0.02	0.11
Verify		0.02	0.17	0.05	-0.03	-0.16	0.17	-0.02	-0.31	0.03	-0.16
PSI- BLAST		1.60E-62	1.10E-29	9.00E-15	1.10E-39	1.10E-24	3.60E-17	1.80E-22	1.60E-21	3.60E-27	0
End		220	322	715	411	319	708	319	305	319	412
Start		21	132	524	234	143	512	139	141	143	21
Chain		Ξ	∢	a	α	V	∢	∢		∢	В
PDB	*	1f4w) f6a	1166f	1fc2	lfcg	1fith	1fbi	1hnf	Ihng	ligt
SEQ	ö	846	846	846	846	846	846	846	846	846	846

PDB annotation	REGION, IMMUNOGLOBULIN	IMMUNOGLOBULIN INTACT IMMUNOGLOBULIN, V REGION, C REGION, HINGE REGION	IMMUNOGLOBULIN INTACT IMMUNOGLOBULIN, V REGION, C REGION, HINGE REGION	COMPLEX (IMMUNOGLOBULIN/RECEPTOR) IMMUNOGLOBULIN FOLD, TRANSMEMBRANE, GLYCOPROTEIN, RECEPTOR, 2 SIGNAL, COMPLEX (IMMUNOGLOBULIN/RECEPTOR)	COMPLEX (IMMUNOGLOBULINRECEPTOR) TCR (VAPIHA VBETA DOMAIN; T-CELL VAPIHALOR, STRAND SWITCH, FAB, ANTICLONOTYPIC, 2 (IMMUNOGLOBULINRECEPTOR)					
Coumpound		IGGI INTACT ANTIBODY MAB61.1.3; CHAIN: A, B, C, D	IGGI INTACT ANTIBODY MAB61.1.3; CHAIN: A, B, C, D	INTERLEUKIN-I BETA; CHAIN: A; TYPE I INTERLEUKIN-I RECEPTOR; CHAIN: B;	KB5-C20 T-CELL ANTIGEN RECEPTOR; CHAIN: A, B; ANTIBODY DESIRE-1; CHAIN: L, H;	IMMUNOGLOBULIN ANTI- PHOSPHATIDYLINOSITOL SPECIFC PHOSPHOLIPASE C DIABODY ILMK 3 SYNONYMS: LSMK 16 DIABODY, SINGLE-CHAIN FV DIMER ILMK 4	IMMUNOGLOBULIN ANTIGEN-BINDING FRAGMENT (FAB) (IGG2B, KAPPA) IMAM 3	IMMUNOGLOBULIN IMMUNOGLOBULIN GI (IGGI) (MCG) WITH A HINGE DELETION IMCO 3	IMMUNOGLOBULIN IMMUNOGLOBULIN GI (IGGI) (MCG) WITH A HINGE DELETION IMCO 3	IMMUNOGLOBULIN IMMUNOGLOBULIN FAB FRAGMENT (MCPC\$603)
SeqFold score		110.16							123.28	
PMF			0.24	0.49	-0.17	0:01 ,	0.22	0.04		90.0
Verify score			-0.05	0.21	0.11	-0.23	-0.05	-0.17		-0.26
PSI- BLAST		3.20E-95	3.20E-95	5.40E-29	1.10E-49	3.20E-39	4.80E-62	1.60E-93	1.60E-93	8.00E-46
End		412	411	503	413	206	220	411	416	210
Start		13	21	225	235	21	21	20	24	21
Chain		æ	В	æ	#	4 .	H	н	H	H
PDB	1	ligy	ligy	1itb	1,465	11mk	Ima m	lmc o	lmc o	J mc
SEQ	ö	846	846	846	846	846	846	846	846	846

PDB annotation		CELL ADHESION PROTEIN CELL ADHESION PROTEIN, RGD, EXTRACELLULAR MATRIX, 2 HEPARIN-BINDING, GLYCOPROTEIN		COMPLEX (INMUNORECEPTOR/INMUNOGLOBU LIN) COMPLEX (INMUNORECEPTOR/INMUNOGLOBU	LIN)	IMMUNOGLOBULIN IMMUNOGLOBULIN,	IMMUNOGLOBULIN VARIABLE HEAVY (VH) DOMAIN, VARIABLE LIGHT (VL) ANTIBODY FRAGMENT, MULTIVALENT ANTIBODY, DIABODY, DOMAIN 2 SWAPPING, IMMUNOGLOBULIN	STRUCTURAL PROTEIN INTEGRIN, HEMIDESMOSOME, FIBRONECTIN, CARCINOMA, STRUCTURAL 2 PROTEIN	STRUCTURAL PROTEIN INTEGRIN, HEMIDESMOSOME, FIBRONECTIN, CARCINOMA, STRUCTURAL 2 PROTEIN	IMMUNE SYSTEM FAB, PORA, NEISSERIA MENINGITIDIS, PORIN	IMMUNOGLOBULIN IMMUNOGLOBULIN, SINGLE-CHAIN FV, ANTI-CARCINOEMBRYONIC 2 ANTIGEN	STRUCTURAL PROTEIN TENASCIN,
Coumpound	1MCP 4	FIBRONECTIN; CHAIN: NULL;	IMMUNOGLOBULIN IGG JEL 103 FAB FRAGMENT COMPLEXED WITH IMRD 3 INOSINE-5:-DIPHOSPHATE IMRD 4	N15 ALPHA-BETA T-CELL RECEPTOR; CHAIN: A, B, C, D; H57 FAB; CHAIN: E, F, G, H		NIG9 (IGGI=LAMBDA=); CHAIN: L, H;	SINGLE-CHAIN ANTIBODY FRAGMENT; CHAIN: A, C;	INTEGRIN BETA-4 SUBUNIT; CHAIN: A, B;	INTEGRIN BETA-4 SUBUNIT; CHAIN: A, B;	ANTIBODY; CHAIN: H, L; PROTEIN G-PRIME; CHAIN: A; MAJOR OUTER MEMBRANE PROTEIN PI.16; CHAIN: P.	MFE-23 RECOMBINANT ANTIBODY FRAGMENT; CHAIN: A;	TENASCIN; CHAIN: A, B;
SeqFold score												
PMF		0.15	-0.06	-0.05		0.13	0.04	0.21	0.24	-0.01	0.25	0.19
Verify score		-0.03	0.05	0.05		0.14	-0.11	0.08	0.1	0.12	0.24	0.04
PSI- BLAST		3.60E-21	9.60E-64	4.80E-62		1.60E-63	3.20E-43	5.40E-15	1.80E-23	3.20E-63	8.00E-40	5.40E-19
End		715	220	220		220	206	599	715	220	211	713
Start		512	21	21		21	21	449	808	21	21	512
Chain			H	r.		н	∢	V	¥	н	V .	Ą
PDB ID		Imfi	1mrd	Infd		lngp	Inqb	барі	lqg3	lqkz	lqok	1qr4
S B SE		846	846	846		846	846	846	846	846	846	846

PDB annotation	FIBRONECTIN TYPE-III, HEPARIN, EXTRACELLULAR 2 MATRIX, ADHESION, FUSION PROTEIN, STRUCTURAL PROTEIN			IMMUNOGLOBULIN ANTI- NITROPHENOL, LAMBDA LIGHT CHAIN, IMMUNOGLOBULIN	CELL ADHESION ICAM-2; IMMUNOGLOBULIN FOLD, CELL ADHESION, GLYCOPROTEIN, 2 TRANSMEMBRANE, REPEAT, SIGNAL	IMMUNE SYSTEM PS8 NATURAL KILLER CELL RECEPTOR; KIR, NATURAL KILLER RECEPTOR, INHIBITORY RECEPTOR, IMMUNOGLOBULIN			A; IMMUNE SYSTEM CD32; RECEPTOR, FC, CD32, IMMUNE SYSTEM	Z	FNDOCYTOSIS/EXOCYTOSIS NSECI:	The contract of the contract o
Coumpound		HUMAN VASCULAR CELL ADHESION MOLECULE-1; 1VCA 4 CHAIN: A, B; 1VCA	E8 ANTIBODY; CHAIN: L, H; CYTOCHROME C; CHAIN: F;	FAB FRAGMENT; CHAIN: NULL;	INTERCELLULAR ADHESION MOLECULE-2; CHAIN: NULL;	MHC CLASS I NK CELL RECEPTOR PRECURSOR; CHAIN: A;	IMMUNOGLOBULIN IG*A FAB FRAGMENT (1539) (GALACTAN-BINDING) 2FBJ	FC GAMMA RIIB; CHAIN: A;	FC GAMMA RIIB; CHAIN: A;	IMMUNOGLOBULIN FAB FRAGMENT FROM HUMAN IMMUNOGLOBULIN IGGI (LAMBDA, HIL) 8FAB 3	SVALTA VINI DINIDING	STALASIN BINDING
SeqFold score										•		_
PMF score		0.74	-0.14	-0.15	0.09	0.09	0.19	0.81	0.21	0	:: 6	= = =
Verify		0.19	60.0	0	0	0.29	0.12	0.13	-0.04	-0.35	3	0:0 0:0
PSI- BLAST		3.60E-24	8.00E-50	8.00E-53	1.80E-34	7.20E-28	3.20E-50	7.20E-36	9.00E-29	9.60E-28		1.60E-09
End		319	413	602	326	320	210	323	410	407		480
Start AA		137	236	419	135	134	21	135	226	234		299
Chain		V	H	H		<	н	4	4	∢		m
PDB		lvca	lwej	lyuh	lzxd	2dli	2fbj	2fcb	2fcb	8fab		1 d
SEQ	ÖZ	846	846	846	846	846	846	846	846	846		847

PDB annotation	MULTI-SUBUNIT	HYDROLASE BETA-ALPHA BARREL	ANTIMICROBIAL PROTEIN DISULFIDE-RICH	HYDROLASE HYDROLASE, CHITIN DEGRADATION	HYDROLASE HYDROLASE, CHITIN DEGRADATION	HYDROLASE BETA-ALPHA (TIM) BARREL	HYDROLASE BETA-ALPHA (TIM) BARREL													-				
Coumpound	SYNTAXIN 1A; CHAIN: B;	CHITINASE 1: CHAIN: A:	TACHYCITIN; CHAIN: A	CHITINASE B; CHAIN: A, B;	CHITINASE B; CHAIN: A, B;	CHITINASE A; CHAIN: A;	CHITINASE A; CHAIN: A;	HYDROLASE (GLUCOSIDASE) ENDO- BETA-N-	ACETYLGLUCOSAMINIDAS E H, ENDO H (E.C.3.2.1.96) 1EDT 3	HYDROLASE(GLUCOSIDASE	ACETYLGLUCOSAMINIDAS	E F1 (E.C.3.2.1.96) 2EBN 3 ·	(ENDOGL YCOSIDASE F1, ENDO F1) 2EBN 4	HYDROLASE(GLUCOSIDASE	ACETYLGLUCOSAMINIDAS	E F1 (E.C.3.2.1.96) 2EBN 3	(ENDOGL YCOSIDASE F1, ENDO F1) 2EBN 4	DNA-BINDING PROTEIN	(HOMEODOMAIN) MUTANT	WITH CYS 39 1AHD 3	REPLACED BY SER (C39S)	COMPLEX WITH DNA (NMR,	IAHD 4 16 STRUCTURES) IAHD 5	DNA-BINDING PROTEIN
SeqFold score																		77.33						
PMF		-	-0.09	_	_	_	_	0.13		90.0				0.77										
Verify score		0.35	0.19	0.47	0.51	0.41	0.33	-0.15		0.25				-0.3										0.13
PSI- BLAST		4.80E-63	1.60E-10	1.10E-81	1.60E-59	3.60E-79	9.60E-70	4.80E-05		7.20E-51				2900'0				7.20E-33						6.40E-29
End AA		297	367	265	566	281	260	146		504				Ξ				351						337
Start AA		2	323	_	2		4	4		I				4				284						286
Chain TD		 	¥	4	∢	4	¥											۵.						_ይ
PDB		1d2k	1dqc	1.00 E+15	1.00 E+15	1edq	ledq	ledt		2ebn				2ebn				lahd						1ahd
SEQ SO SO SO SO SO SO SO SO SO SO SO SO SO		848	848	848	848	848	848	848		848				848				849						849

PDB annotation			PROTEIN/DNA HOMEODOMAIN, DNA, COMPLEX, DNA-BINDING PROTEIN, PROTEIN/DNA	PROTEIN/DNA HOMEODOMAIN, DNA, COMPLEX, DNA-BINDING PROTEIN, PROTEIN/DNA	TRANSCRIPTION/DNA ULTRABITHORAX; PBX PROTEIN; DNA BINDING, HOMEODOMAIN, HOMEOTIC PROTEINS, DEVELOPMENT, 2 SPECIFICITY	TRANSCRIPTION/DNA ULTRABITHORAX; PBX PROTEIN; DNA BINDING, HOMEODOMAIN, HOMEOTIC PROTEINS, DEVELOPMENT, 2 SPECIFICITY	TRANSFERASE DINUCLEOTIDE- BINDING MOTIF, PHOSPHORIBOSYL TRANSFERASE	
Coumpound	ANTENNAPEDIA PROTEIN (HOMEODOMAIN) MUTANT WITH CYS 39 1 AHID 3 REPLACED BY SER (C39S) COMPLEX WITH DNA (NMR, 1 AHID 4 16 STRUCTURES)	DNA-BINDING PROTEIN ANTENNAPEDIA PROTEIN (HOMEODOMAIN) MUTANT WITH CYS 39 1AHD 3 REPLACED BY SER (C39S) COMPLEX WITH DNA (NMR, 1AHD 4 16 STRUCTURES)	HOMEOBOX PROTEIN HOX- B1; CHAIN: A; PBX1; CHAIN: B; DNA CHAIN: D; DNA CHAIN: E;	HOMEOBOX PROTEIN HOX- B1; CHAIN: A; PBX1; CHAIN: B; DNA CHAIN: D; DNA CHAIN: E;	ULTRABITHORAX HOMEOTIC PROTEIN IV; CHAIN: A; HOMEOBOX PROTEIN EXTRADENTICLE; CHAIN: B; DNA (5'- CHAIN: C; DNA (5'- CHAIN: D;	ULTRABITHORAX HOMEOTIC PROTEIN IV; CHANI: A; HOMEOBOX PROTEIN EXTRADENTICLE; CHANI: B; DNA (5'- CHAIN: C; DNA (5'- CHAIN: D;	NICOTINATE MONONUCLEOTIDE:5,6- CHAIN: A;	DNA-BINDING FUSHI TARAZU PROTEIN (HOMEODOMAIN) (NMR, 20
SeqFold score			62.11		65.54			68.94
PMF		-		1		66.0	-0.19	
Verify score		-0.38		-0.08		-0.11	0.15	
PSI- BLAST	1	7.20E-33	5.40E-32	5.40E-32	5.40E-31	5.40E-31	7.20E-11	1.30E-27
End		351	346	346	342	342	238	351
Start AA		286	274	286	285	286	35	283
Chain ID		D.	V	∢	<	∢	¥	
PDB ID		lahd	1672	1672	168i	1881	140s	1ftz
SEQ NO.		849	849	849	849	849	849	849

PDB annotation			•					•		·	COMPLEX (DNA-BINDING PROTEIN/DNA) HD; HOMEODOMAIN, COMPLEX (DNA-BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) HD; HOMEODOMAIN, COMPLEX (DNA-BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING
Coumpound	STRUCTURES) 1FTZ 3	DNA-BINDING FUSHI TARAZU PROTEIN (HOMEODOMAIN) (NMR, 20 STRUCTURES) IFTZ 3	DNA-BINDING PROTEIN OCT-1 (POU DOMAIN) 10CT 3	DNA-BINDING PROTEIN ANTENNAPEDIA PROTEIN (HOMEODOMAIN) MUTANT WITH CYS 39 ISAN 3 REPLACED BY SER AND RESIDUES 1-6 DELETED	(C39S,DEL 1-6) ISAN 4 (NMR, 20 STRUCTURES) ISAN 5	DNA-BINDING PROTEIN ANTENNAPEDIA PROTEIN (HOMEODOMAIN) MUTANT	WITH CYS 39 ISAN 3 REPLACED BY SER AND RESIDUES I-6 DELETED	(C39S,DEL 1-6) ISAN 4 (NMR, 20 STRUCTURES) ISAN 5	DNA-BINDING PROTEIN ANTENNAPEDIA PROTEIN (HOMEODOMAIN) MUTANT WITH CYS 39 15AN 3	REPLACED BY SER AND RESIDUES 1-6 DELETED (C39S, DEL 1-6) ISAN 4 (MMR, 20 STRUCTURES) ISAN 5	ANTENNAPEDIA PROTEIN; CHAIN: A, B; DNA; CHAIN: C, D, E, F;	ANTENNAPEDIA PROTEIN; CHAIN: A, B; DNA; CHAIN: C, D, E, F;	ANTENNAPEDIA PROTEIN;
SeqFold score				72.13								90.69	
PMF		0.84	0.12						-		-		_
Verify score		-0.31	-0.2			-0.21			-0.2		0.11		0.27
PSI- BLAST		1.30E-27	7.20E-34	5.40E-30		1.40E-26			5.40E-30		1.30E-27	9.00E-30	9.00E-30
End		337	344	351		337			351		337	344	344
Start AA		284	205	290		291			292		289	289	289
Chain ID			U								٧	⋖	
PDB UD		2 1	- ö	Isan		Isan			1san		9ant	9ant	9ant
SEQ EQ		849	849	849		849			849		849	849	849

		 		_						_	_			_						_					_
PDB annotation	PROTEIN/DNA) HD; HOMEODOMAIN, COMPLEX (DNA-BINDING PROTEIN/DNA)	COMPLEX (TRANSMEMBRANE/GLYCOPROTEIN) MHC GLYCOPROTEIN, COMPLEX (TRANSMEMBRANE/GLYCOPROTEIN)	COMPLEX (TRANSMEMBRANE/GLYCOPROTEIN) MHC GLYCOPROTEIN, COMPLEX (TRANSMEMBRANE/GLYCOPROTEIN)	COMPLEX (MHC PROTEIN/ANTIGEN)	DRA, DRB1 01010; COMPLEX (MHC PROTEIN/ANTIGEN).	HISTOCOMPATIBILITY ANTIGEN	COMPLEX (MHC PROTEIN/ANTIGEN)	DRA, DRB1 01010; COMPLEX (MHC	PROTEIN/ANTIGEN, HISTOCOMPATIBILITY ANTIGEN		IMMUNE SYSTEM HLA-DR2, MYELIN	BASIC PROTEIN, MULTIPLE	SCLEROSIS, 2 AUTOIMMUNITY, IMMUNE SYSTEM	IMMUNE SYSTEM HLA-DR2, MYELIN	BASIC PROTEIN, MULTIPLE	SCLEROSIS, 2 AUTOIMMUNITY, IMMUNE SYSTEM	IMMUNE SYSTEM HLA-DR4; HLA-	DR4; SEB, SUPERANTIGEN; COMPLEX	IMMUNE SYSTEM					IMMUNE SYSTEM MHC CLASS II	DR2A
Coumpound	CHAIN: A, B; DNA; CHAIN: C, D, E, F;	HLA-DR3; CHAIN: A, B; CLIP; CHAIN: C;	HLA-DR3; CHAIN: A, B; CLIP; CHAIN: C;	HLA-DRI CLASS II	HISTOCOMPATIBILITY PROTEIN: CHAIN: A B D E	G, H, I, K; HLA-A2; CHAIN: C, F. I. L:	HLA-DRI CLASS II	HISTOCOMPATIBILITY	G H I K: HLA-A2: CHAIN: C.	F. I. L.	HLA-DR2; CHAIN: A, D; HLA-	DR2; CHAIN: B, E; HLA-DR2;	CHAIN: C, F;	HLA-DR2; CHAIN: A, D; HLA-	DR2; CHAIN: B, E; HLA-DR2;	CHAIN: C, F;	HLA CLASS II	HISTOCOMPATIBILITY	CLASS II	HISTOCOMPATIBILITY	ANTIGEN; CHAIN: B;	ENTEROTOXIN TYPE B;	CHAIN: C; PEPTIDE NHIBITOR: CHAIN: D:	MAJOR	HISTOCOMPATIBILITY
SeqFold score		54.43					55.9							52.4											
PMF score			0.93	0.63							0.87						0.92							0.82	
Verify score			-0.32	-0.33							-0.22						-0.33							-0.31	
PSI- BLAST		3.20E-66	3.20E-66	6.40E-69			6.40E-69				1.60E-69			1.60E-69			4.80E-62							3.20E-67	
End AA		157	147	147			157				147			157			147							147	
Start AA		33	34	33			33				32			32	!		31							30	
Chain		æ	B	В			В				В			B	١_		В							В	
PDB CD		1a6a	1a6a	land	•		lagd				1bx2			1bx2			1d5	E						1441	
SEQ B	Š	851	851	851			851				851			851	:		851		_					821	

PDB annotation	AT A I VALUE DATE TO A DESCRIPTION OF THE PERSON OF THE PE	HISTOCOMPATIBILITY ANTIGEN, HISTOCOMPATIBILITY ANTIGEN, MHC, PEPTIDE COMPLEX	OVALBUMIN PEPTIDE	MHC II MHC II, CLASS II MHC, 1-4, OVALBUMIN PEPTIDE	HISTOCOMPATIBILITY ANTIGEN HISTOCOMPATIBILITY ANTIGEN	HISTOCOMPATIBILITY ANTIGEN HISTOCOMPATIBILITY ANTIGEN	HISTOCOMPATIBILITY ANTIGEN HISTOCOMPATIBILITY ANTIGEN	MHC II MHC II, CLASS II MHC I-AD	MHC II MHC II, CLASS II MHC I-AD	BLOOD COAGULATION BLOOD COAGULATION, EGF, HYDROLASE, SERINE PROTEASE	STRUCTURAL PROTEIN INTEGRIN- BINDING PROTEIN, INV GENE		BLOOD COAGULATION, SERINE PROTEASE, COMPLEX, CO-FACTOR, 2 RECEPTOR ENZYME, INHIBITOR, GLA, EGF, 3 COMPLEX (SERINE PROTEASE/COFACTOR/LIGAND)	
Coumpound	COMPLEX ALPHA CHAIN; CHAIN: A, D; MAJOR HISTOCOMPATIBILITY COMPLEX BETA CHAIN; CHAIN: B, E; MYELIN BASIC PROTEIN; CHAIN: C, F;	MHC CLASS II I-AK; CHAIN: A, B, P; HEN EGGWHITE LYSOZYME PEPTIDE	MHC CLASS II I-AD; CHAIN: A, B;	MHC CLASS II I-AD; CHAIN: A, B;	MHC CLASS II I-EK; CHAIN: A, B, C, D;	MHC CLASS II I-EK; CHAIN: A, B, C, D;	MHC CLASS II I-EK; CHAIN: A, B. C, D;	MHC CLASS II I-AD; CHAIN: A, B;	MHC CLASS II I-AD; CHAIN: A, B;	FACTOR VII; CHAIN: NULL;	INVASIN; CHAIN: A;	GLYCOSYLTRANSFERASE CYCLODEXTRUN GLUCANOTRANSFERASE (E.C.2.4.1.19) (CGTASE) 1CYG 3	BLOOD COAGULATION FACTOR VIIA; CHAIN: L, H; SOLUBLE TISSUE FACTOR; CHAIN: T, U; D-PHE-PHE- ARG-	(DFFRCMK) WITH CHAIN: C;
SeqFold score			52.61			50.29		52.73				·		
PMF		0.46		0.71	6.0		0.87		-	0.17	-0.19	-0.19	0.04	
Verify		-0.4		-0.39	-0.27		-0.43		-0.5	-0.42	0.05	0.07	-0.24	
PSI- BLAST		4.80E-53	9.60E-57	9.60E-57	1.40E-63	1.40E-63	8.00E-64	3.20E-58	3.20E-58	0.0072	3.60E-19	7.20E-15	0.009	
End		147	157	147	147	157	147	157	147	474	229	226	475	
Start		36	23	35	29	∞	24	22	26	443	2	17	404	
Chain		В	В	В	Д	В	В	В	В		A		<u>ا</u>	
PDB		liak	liao	liao	liea	lica	1 jeb	2iad	2iad	1649	lcwv	lcyg	ldan	
SEQ Second	2	851	851	851	851	851	851	158	851	856	856	958	856	

PDB annotation	HYDROLASEHYDROLASE NHIBITOR PROTEIN-PEPTIDE COMPLEX	HYDROLASE NEURAMINIDASE; HYDROLASE, GLYCOSIDASE	BLOOD CLOTTING FACTOR VII, BLOOD COAGULATION, EGF-LIKE DOMAIN, BLOOD 2 CLOTTING	GROWTH FACTOR NEU DIFFERENTIATION FACTOR (RAT), ACETYLCHOLINE GROWTH FACTOR		GLYCOSYLTRANSFERASE TRANSFERASE, GLYCOSYLTRANSFERASE, CALCIUM, SIGNAL	HYDROLASE HYDROLASE, LIPID DEGRADATION, PANCREATIC LIPASE	HYDROLASE HYDROLASE, LIPID DEGRADATION, PANCREATIC LIPASE	COMPLEX (HYDROLASE/COFACTOR) TRIACYLGLYCEROL LIPASE; COMPLEX (HYDROLASE/COFACTOR), LIPID DEGRADATION	COMPLEX (HYDROLASE/COFACTOR) TRIACYLGLYCEROL LIPASE; COMPLEX (HYDROLASE/COFACTOR), LIPID DEGRADATION
Coumpound	DES-GLA FACTOR VIIA (HEAVY CHAIN); CHAIN: H, I; DES-GLA FACTOR VIIA (LIGHT CHAIN); CHAIN: L, M; (DPN)-PHE-ARG; CHAIN: C, D; PEPTIDE E-76; CHAIN: X, Y;	SIALIDASE; CHAIN: NULL:	BLOOD COAGULATION FACTOR VII; CHAIN: A;	HEREGULIN-ALPHA; CHAIN: NULL;	GROWTH FACTOR HEREGULIN-ALPHA (EPIDERMAL GROWTH FACTOR-LIKE DOMAIN) IHRE 3 (NMR, MINIMIZED AVERAGE STRUCTURE) IHRE 4	CYCLODEXTRIN GLUCANOTRANSFERASE; CHAIN: A, B;	PANCREATIC LIPASE RELATED PROTEIN 2; CHAIN: A;	PANCREATIC LIPASE RELATED PROTEIN 2; CHAIN: A;	TRIACYLGLYCEROL ACYL- HYDROLASE; CHAIN: A, C; COLIPASE; CHAIN: B, D	TRIACYLGLYCEROL ACYL- HYDROLASE; CHAIN: A, C; COLIPASE; CHAIN: B, D
SeqFold score									167.54	
PMF	0.28	-0.19	0.11	0	0.23	-0.19	_	-		-
Verify	0.07	0.22	-0.28	-0.21	-0.7	0.04	0.85	0.81		0.65
PSI- BLAST	0.0036	7.20E-13	0.0036	0.0072	0.0072	5.40E-12	3.60E-83	4.80E-65	1.60E-66	1.60E-66
End AA	472	232	475	482	254	171	256	276	276	276
Start	429	18	443	429	230	-	18	<u>8</u>	82	18
Chain	دا		4			4	4	<	∢ .	¥.
PDB DD	ldva	lent	1f7e	1hae	1hrc	lpa m	15u8	1bu8	1eth	1eth
SEQ EQ	856	856	856	856	856	826	857	857	857	857

					··		
PDB annotation	SERINE ESTERASE RELATED PROTEIN 2 LIPASE; SERINE ESTERASE, HYDROLASE, LIPID DEGRADATION, PANCREAS, 2 GLYCOPROTEIN, CHIMERIC	SERINE ESTERASE RELATED PROTEIN 2 LIPASE, SERINE STERASE, HYDROLASE, LIPID DEGRADATION, PANCREAS, 2 GLYCOPROTEIN, CHIMERIC	SERINE ESTERASE RELATED PROTEIN 2 LIPASE, SERINE SETERASE. HYDROLASE, LIPID DEGRADATION, PANCREAS, 2 GLYCOPROTEIN, CHIMERIC				
Coumpound	RP2 LIPASE; CHAIN: NULL;	RP2 LIPASE; CHAIN: NULL;	RP2 LIPASE; CHAIN: NULL;	HYDROLASE(CARBOXYLIC ESTERASE) LIPASE (EC.3.1.1.3) (TRIACYLGLYCEROL HYDROLASE) 1HPL 3	HYDROLASE(CARBOXYLIC ESTERASE) LIPASE (E.C.3.1.1.3) COMPLEXED WITH COLIPASE AND INHIBITED ILPB 3 BY UNDECANE PHOSPHONATE METHYL ESTER (TWO CONFORMATIONS) 11.PB 4	HYDROLASE(CARBOXYLIC ESTERASE) LIPASE (E.C.3.1.1.3) COMPLEXED WITH COLIPASE AND INHIBITED 1LPB 3 BY UNDECANE PHOSPHONATE METHYL ESTER (TWO CONFORMATIONS) 1LPB 4	HYDROLASE(CARBOXYLIC ESTERASE) LIPASE (E.C.3.1.1.3) COMPLEXED WITH COLIPASE AND INHIBITED 1LPB 3 BY
SeqFold score	176.81			167		173.41	
PMF		1	-		-		-
Verify score		0.87	0.76		0.83		0.81
PSI- BLAST	1.80E-80	1.80E-80	3.20E-65	1.60E-76	1.10E-79	1.10E-79	1.60E-65
End AA	276	256	276	274	256	276	276
Start AA	18	61	61	81	8	81	18
Chain ID				. ₹	æ	m.	æ
PDB	1gpl	1gpl	lgpl	1hpl	11pb	11pb	qdll
SEQ B B	857	857	857	857	857	857	857

S	Start	End	PSI- BLAST	Verify score	PMF	SeqFold	Coumpound	PDB annotation
							UNDECANE PHOSPHONATE METHYL ESTER (TWO CONFORMATIONS) ILPB 4	
18 257	257		5.40E-83	0.69	-		PANCREATIC LIPASE RELATED PROTEIN 1; CHAIN: NULL:	HYDROLASE HYDROLASE, LIPID DEGRADATION, PANCREATIC LIPASE
18 274	274	T.	5.40E-83			183.05	PANCREATIC LIPASE RELATED PROTEIN 1; CHAIN: NULL;	HYDROLASE HYDROLASE, LIPID DEGRADATION, PANCREATIC LIPASE
18 276	276		1.60E-63	0.71	_		PANCREATIC LIPASE RELATED PROTEIN 1; CHAIN: NULL;	HYDROLASE HYDROLASE, LIPID DEGRADATION, PANCREATIC LIPASE
-	ļ	Γ						
14 156	156		0.00036	-0.08	0		SYNTAXIN BINDING PROTEIN 1; CHAIN: A; SYNTAXIN 1A; CHAIN: B;	ENDOCYTOSIS/EXOCYTOSIS NSEC1; PROTEIN-PROTEIN COMPLEX, MULTI-SUBUNIT
151 308	308	I	1.30E-44	0.28	0.07		TRANSCRIPTIONAL REPRESSOR TUP1; CHAIN: A, B. C.	TRANSCRIPTION INHIBITOR BETA- PROPELLER
132 298	562		8.00E-46	0.19	0.12		GT-ALPHA/GI-ALPHA CHIMERA; CHAIN: A; GT- BETA; CHAIN: B; GT- GAMMA; CHAIN: G;	COMPLEX (GTP-BINDING/TRANSDUCER) BETA1, TRANSDUCIN BETA SUBUNIT; GAMMA1, TRANSDUCIN GAMMA SUBUNIT; COMPLEX (GTP-BINDING/TRANSDUCER), G PROTEIN, HETEROTRIMER 2 SIGNAL TRANSDUCTION
300	308		1.10E-40			52.34	GT-ALPHA/GI-ALPHA CHIMERA; CHAIN: A; GT- BETA; CHAIN: B; GT- GAMMA; CHAIN: G;	COMPLEX (GTP-BINDING/TRANSDUCER) BETA!, BINDING/TRANSDUCER) BETA!, TRANSDUCIN BETA SUBUNIT; GAMMAI, TRANSDUCIN GAMMA SUBUNIT; COMPLEX (GTP-BINDING/TRANSDUCER), HETEROTRIMER 2 SIGNAL TRANSDUCTION
95 309	308	_	1.10E-40	0.13	-0.11		GT-ALPHA/GI-ALPHA CHIMERA; CHAIN: A; GT- BETA; CHAIN: B; GT- GAMMA; CHAIN: G;	COMPLEX (GTP- BINDINGTRANSDUCER) BETA1, TRANSDUCIN BETA SUBUNIT; GAMMA1, TRANSDUCIN GAMMA SUBUNIT; COMPLEX (GTP- BINDINGTRANSDUCER), G PROTEIN,

			Т	·T			1	_		٦	_		_							_					Γ	
PDB annotation	HETEROTRIMER 2 SIGNAL TRANSDUCTION	OXIDOREDUCTASE ENZYME, NITRITE REDUCTASE, OXIDOREDUCTASE, DENITRIFICATION, 2 ELECTRON TRANSPORT, PERPLASMIC		COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN		COMPLEX (ZINC FINGER/DNA)	FINGER, DNA-BINDING PROTEIN		COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN		COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN		COMPLEX (ZINC FINGER/DNA) ZINC	INTERACTION PROTEIN DESIGN 2	CRYSTAL STRUCTURE, COMPLEX	(ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC	NITER ACTION PROTEIN DESIGN 2	CRYSTAL STRIPTIBE COMPLEX	(ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC	FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2
Coumpound		CYTOCHROME CDI NITRITE REDUCTASE, CHAIN: A, B;		QGSR ZINC FINGER PEPTIDE, CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	OGSR ZINC FINGER	PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE	BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER	OLIGONUCLEOTIDE	BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER	PEPTIDE; CHAIN: A; DUPLEX OI.IGONIICI EOTIDE.	BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER	PEPTIDE; CHAIN: A; DUPLEX	BINDING SITE; CHAIN: B, C;	DNA; CHAIN: A, B, D, E;	PROTEIN CHAIN: DE G.	Indicat, Charle 5, 1, 5,		DNA; CHAIN: A, B, D, E;	DEOTERNOOS ZINC FINGER	The leaf, clining c, t, c.		DNA; CHAIN: A, B. D, E;	CONSENSUS ZINC FINGER PROTEIN: CHAIN: C, F, G;
SeqFold score																										
PMF score		-0.12		0.15				-			96.0			0.99			0.07				0.52				_	
Verify score		0		-0.3	9			0.48			0.27			0.41			-0.35				-0.16				0.1	
PSI- BLAST		1.30E-14		1.60E-25	1.80E-39			9.00E-38			3.60E-39			6.40E-31			1.10E-39				3.20E-43				1.40E-44	
End		304		221	277			306			640			940			193				221				249	
Start AA		221		141	197			225			260			260			112				140				168	
Chain ID		¥		∢	4			Ą			A			4			O				ပ				U	
PDB ID		1qks		laih	lalh			lalh			lalh			lalh			1mc	>			lmc	>			1me	у
SEQ NO.		829		860	860			860			860			98			860				860				860	

PDB annotation	CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGERDNA) ZINC FINGER, PROTEN-DNA INTERACTION, PROTEN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGERDNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGERDNA)	COMPLEX (ZINC FINGENDINA) ZINC FUNGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGERDINA) ZINC FINGER, PROTEIN-DNA FINGERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGERDINA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGENDINA) ZINC MTROBE, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGERDNA)	COMPLEX (ZINC FINGENDINA) ZINC MIGGE, PROTEIN-DING PITERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGERDINA)	COMPLEX (ZINC FINGERODNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX
Coumpound	CR	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G; CR	# ::	Ж.:s	E ::	E ::	.; ER	DNA; CHAIN: A. B. D. E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G; C	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;
SeqFold score									
PMF		-	-	<u></u>	-			-	
Verify		0.23	0.55	0.49	0.62	0.34	0.27	0.42	0.16
PSI- BLAST		1.60E-46	1.60E-47	1.40E-48	3.20E-49	8.00E-50	1.60E-50	1.60E-50	4.80E-50
End		277	305	333	361	389	417	445	473
Start		196	224			308	336	364	392
Chain		U	O	U	v	U	U	U	U
PDB UD		y y	Ine y	√ ne	lme y	Ime	1me y	Jme y) y
SEQ ID	ÖŽ	860	98	860	860	860	860	098	860

PDB annotation	(ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGERDNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA COMPLEX CTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA FINERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL, STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
Coumpound		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;
SeqFold score							101.87		
PMF score		0.98		96:0	-	1		-	-
Verify score		-0.22	0.1	0.15	0.43	0.69		0.71	0.39
PSI- BLAST		7.20E-33	9.60E-50	3.20E-47	3.20E-47	9.60E-50	8.00E-50	8.00E-50	1.60E-49
End		200	501	528	556	584	585	612	640
Start		392	420	448	476	503	503	531	559
Chain ID		U	U	U	O	U	U	ပ	ပ
PDB	1	y y	Jme y	ime y	Jme y	Jme y	y y	y Jac	lme y
SEQ D		098	988	098	860	860	9860		860

					104	17.	DAKE	SooFold	Coumpound	PDB annotation
SEQ EQ	EDB EDB		Start	¥ A	BLAST	score	score	score		
.00 860	γ γ	D D	306	333	3.60E-14	0.06	0.94		DNA; CHAN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX ZINC FINGER/DNA)
860	Ime y	O	306	333	4.80E-13	0.06	0.94		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
860	8	<	113	258	4.80E-31	-0.25	0.12		TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	COMPLEX (TRANSCRYTION REGULATION/DNA) COMPLEX (TRANSCRPTION REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN
860	1466	4	169	314	1.60E-35	0.11	0.99		TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FRIGER PROTEIN
098	148	4	224	393	1.80E-66			105.92	TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	COMPLEX COMPLEX EGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN INITIATION ANSCRIPTION
860	186	<	225	370	4.80E-37	0.21	1		TFIIA; CHAIN: A, D; 3S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	COMPLEX (TRANSCRIPTION COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN COMPLEX COM
860	148	<	281	426	1.60E-38	0.32	-		TFIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN
860	146	4	393	537	1.30E-36	0.16	0.99		TFIIIA; CHAIN: A, D; 5S	COMPLEX (TRANSCRIPTION

PDB annotation	REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION TRANSCRIPTION POLYMERASE III, 2 TRANSCRIPTION NITTATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATIONDNA) YING-YANG 1; TRANSCRIPTION INITIATIOR BLEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATIONDNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2
Coumpound	RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	TFIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YY I; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;
SeqFold							
PMF		0.64	0.68	-	0	0.83	69:0
Verify score		0.11	-0.1	0.06	-0.43	-0.19	-0.37
PSI- BLAST		8.00E-38	6.40E-35	9.60E-35	9.60E-29	1.30E-30	1.80E-34
End		565	598	626	221	249	277
Start		421	449	477		148	150
Chain		V	<	∢	ပ	U	ပ
PDB UD		146	超	146	Iubd	lubd	1ubd
SEQ		860	9860	860	860	860	860

							
PDB annotation	FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (IKANSCAPTION REGULATION/DAN) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATIONDNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATIONDNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATIONDNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER, PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATIONDNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG I; TRANSCRIPTION INITIATION,
Coumpound		YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B:	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA;
SeqFold							
PMF		0.93	0.96	-	-	0.88	
Verify score		-0.19	-0.07	0.19	0.27	0.04	0.2
PSI- BLAST		3.20E-32	1.80E-48	4.80E-34	3.20E-34	5.40E-52	3.20E-34
End		777	333	333	361	445	417
Start		170		227	260	306	316
Chain		U	υ .	O	U	ပ	U
PDB UD	-	pqnı	1 ubd	lubd	lubd	lubd	lubd
SEQ	ÖZ	098	860	860	098	860	860

PDB annotation	INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG I; TRANSCRIPTION INTIATION, INTIATOR ELEMENT, YYI, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DINA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG I; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YYI, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1;
Coumpound	CHADì: A, B;	YYI, CHAIN: C; ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS P5
SeqFold score							
PMF score		–	0.53	0.78	0.96		1
Verify score		-0.14	-0.42	0	-0.07	0.23	0.22
PSI- BLAST		4.80E-35	3.60E-42	6.40E-35	3.20E-31	1.80E-47	3.20E-36
End		473	556	· 102	226	584	584
Start AA		372	390	400	456	481	484
Chain TD		U	U	ပ	ပ	U	၁
PDB U		lubd	. pqn[lubd	Iubd	lubd	lubd
SEQ S B S		098	098	098	098	860	860

PDB annotation	TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG I; TRANSCRIPTION IN/ITIATION, IN/ITIATOR ELEMENT, YYI, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATIONDNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATIONDNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	ZINC FINGER DNA BINDING DOMAIN DNA BINDING MOTIF, ZINC FINGER DNA BINDING DOMAIN	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
Coumpound	NITIATOR ELEMENT DNA; CHAIN: A, B;	YY I; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	SWIS; CHAIN: NULL;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;
SeqFold score			86.51					
PMF		0.98		-	0.72	96:0		-
Verify score		0.43		90.0-	0.15	-0.04	-0.21	-0.1
PSI- BLAST		5.40E-51	5.40E-51	3.20E-34	0.00016	3.20E-31	3.60E-52	5.40E-63
End		640	120	95	642	276	307	363
Start AA		529		539	615	140	170	197
Chain 13		ပ	U	O		∢	∢	∢ _
PDB U		lubd	lubd	lubd	1zfd	2gli	2gli	2gli
SEQ B S		098	. 098	860	860	860	860	860

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PDB annotation	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEINDNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA-BINDING PROTEIN/DNA)	COMPLEX (DNA-BNDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEINDNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING
Coumpound	ZINC FINGER PROTEIN GLJI; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A: DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D:	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII;
SeqFold score								÷			91.33
PMF		86.0	66.0	0.19	0.99	0.49	0.19	0.99	-	-	
Verify score	0.06	0.13	0.25	-0.4	0.03	-0.21	-0.07	-0.02	0.27	0.53	
PSI- BLAST	3.60E-64	1.60E-33	1.80E-64	3.60E-57	1.30E-34	1.60E-32	5.40E-60	1.30E-30	1.40E-35	3.60E-64	3.60E-64
End	391	360	447	558	444	527	614	586	611	640	642
Start	224	722	280	308	316	400	421	456	484	503	503
Chain	¥.	A	V	V V	¥	«	< _	◀	4	∢	
PDB ID	2gli	2gli	2gli	2gli	2gli	2gli	2gli	2gli	2gli	2gli	2gli
SEQ	.098 860	860	860	860	860	860	860	860	098	860	860

							_			_				
PDB annotation	PROTEINIONA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEINIONA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	HYDROL YTIC ENZYME DLH;	DIENELACTONE HYDROLASE, AROMATIC HYDROCARBON CATABOLISM, 2 SERINE ESTERASE, CARBOXYMETHYLENEBUTENOLIDA SE, 3 HYDROLYTIC ENZYME	Mar Old is common and in	COMPLEX OF 1 WO ELONGATION FACTORS EF-TU; EF-TS; ELONGATION FACTOR, NUCLEOTIDE EXCHANGE, GTP-BINDING, 2 COMPLEX OF TWO ELONGATION FACTORS	COMPLEX (TWO ELONGATION	FACTORS) ELONGATION FACTOR FOR TRANSFER, HEAT UNSTABLE, ELONGATION FACTOR FOR TRANSFER, HEAT STABLE.	ELONGATION FACTOR, COMPLEX (TWO ELONGATION FACTORS)		DNA-BINDING HMGA DNA-BINDING HMG-BOX DOMAIN A OF RAT HMGI; IAAB 8 HMG-BOX IAAB 20	DNA-BINDING HMGA DNA-BINDING HMG-BOX DOMAIN A OF RAT HMG1; IAAB 8 HMG-BOX IAAB 20	DNA-BINDING HMGA DNA-BINDING HMG-BOX DOMAIN A OF RAT HMG1; 1AAB 8 HMG-BOX 1AAB 20	DNA BINDING PROTEIN HMG BOX, DNA BENDING, DNA RECOGNITION, CHROMATIN, NMR, DNA 2 BINDING
Coumpound	CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GL11; CHAIN: A; DNA; CHAIN: C, D;	PIENEI ACTONE	DIENELACI ONE HYDROLASE; CHAIN: NULL;		ELONGATION FACTOR TU; CHAIN: A, B, E, F; ELONGATION FACTOR TS; CHAIN: C, D, G, H;	ET ONG ATTON FACTOR TITE	CHAIN: A, C; ELONGATION FACTOR TS; CHAIN: B, D;			HIGH MOBILITY GROUP PROTEIN; 1AAB 5 CHAIN: NULL; 1AAB 6	HIGH MOBILITY GROUP PROTEIN; 1AAB 5 CHAIN: MILL: 1AAB 6	HIGH MOBILITY GROUP PROTEIN: 1AAB 5 CHAIN: NULL: 1AAB 6	NON HISTONE PROTEIN 6 A; CHAIN: A;
SeqFold														
PMF		0.46	200	96.0		99.0	32.0	c/.i			0.89		0.33	0.94
Verify		-0.16		0.02		0.02	١	9 		L	-0.09	99.0	0.41	-0.12
PSI- BLAST		1.40E-36		0.00013		0.00018		0.00018			9.00E-18	5.40E-16	1.80E-06	1.80E-16
End		279		338		2676		2673			185	433	290	180
Start		95		255		2646		2646			106	363	528	112
Chain		A				U		m						4
PDB		2gli		nib1	1	laip		lefu			1aab	1aab	laab	1cg7
SEQ ID	-	988		862		864		864			865	865	865	865

PDB annotation	PROTEIN	DNA BINDING PROTEIN HMG BOX, DNA BENDING, DNA RECOGNITION, CHROMATIN, NMR, DNA 2 BINDING PROTEIN	DNA BINDING PROTEIN HMG BOX, DNA BENDING, DNA RECOGNITION, CHROMATIN, NMR, DNA 2 BINDING PROTEIN	DNA BINDING PROTEIN HMG BOX, DNA BENDING, DNA RECOGNITION, CHROMATIN, NMR, DNA 2 BINDING PROTEIN	DNA BINDING PROTEIN HMG BOX, DNA BENDING. DNA RECOGNITION, CHROMATIN, NMR, DNA 2 BINDING PROTEIN	GENE REGULATION/DNA HMG-1, AMPHOTERN, HEPARIN-BINDING PROTEIN P30; HIGH-MOBILITY GROUP DOMAIN, BENT DNA, PROTEIN-DRUG-DNA 2 COMPLEX, GENE REGULATION/DNA	GENE REGULATION/DNA HMG-1, AMPHOTERIN, HEPARIN-BINDING PROTEIN P30; HIGH-MOBILLTY GROUP DOMAIN, BENT DNA, PROTEIN-DRUG-DNA 2 COMPLEX, GENE REGULATION/DNA	GENE REGULATION/DNA HMG-1, AMPHOTERIN, HEPARIN-BINDING PROTEIN P30; HICH-MOBILITY GROUP DOMAIN, BENT DNA, PROTEIN-DRUG-DNA 2 COMPLEX, GENE REGULATION/DNA	GENE REGULATION/DNA HMG-1, AMPHOTERN, HEPARIN-BINDING PROTEIN P30; HIGH-MOBILITY GROUP DOMAIN, BENT DNA, PROTEIN-DRUG-DNA 2 COMPLEX, GENE REGULATION/DNA
Coumpound		NON HISTONE PROTEIN 6 A; CHAIN: A;	NON HISTONE PROTEIN 6 A; CHAIN: A;	NON HISTONE PROTEIN 6 A; CHAIN: A;	NON HISTONE PROTEIN 6 A; CHAIN: A;	HIGH MOBILITY GROUP 1 PROTEIN; CHAIN: A; DNA (5- D(*CP*CP*(IDO) CHAIN: B; DNA (5- CHAIN: C;	HIGH MOBILITY GROUP 1 PROTEIN; CHAIN: A; DNA (5'- D(*CP*CP*(IDO) CHAIN: B; DNA (5'- CHAIN: C;	HIGH MOBILITY GROUP 1 PROTEIN; CHAIN: A; DNA (5'- D(*CP*CP*(IDO) CHAIN: B; DNA (5'- CHAIN: C;	HIGH MOBILITY GROUP 1 PROTEIN; CHAIN; A; DNA (5'- D(*CP*CP*CIDO) CHAIN; B; DNA (5'- CHAIN; C;
SegFold score									
PMF score		0.83	-	0.42	0.24	0:92	0.71	66:0	0.11
Verify score		0.3	0.45	-0.4	0.28	-0.29	0.02	0.4	0.03
PSI- BLAST		3.60E-07	5.40E-16	7.20E-06	1.80E-06	5.40E-14	3.60E-07	3.60E-15	7.20E-05
End		320	433	516	109	177	320	433	510
Start AA		262	369	476	532	112	262	369	476
Chain		Y	A	V	∢	4	4	4	4
PDB ID		lcg7	lcg7	lcg7	lcg7	1ckt	1ckt	1ckt	lckt
SEQ US		865	865	865	865		865	865	865

								
PDB annotation	GENE REGULATION/DNA HMG-1, AMPHOTERIN, HEPARIN-BINDING PROTEIN P30; HIGH-MOBILLITY GROUP DOMAAIN, BENT DNA, PROTEIN-DRUG-DNA 2 COMPLEX, GENE REGULATION/DNA	STRUCTURAL PROTEIN TWO REPEATS OF SPECTRIN, ALPHA HELICAL LINKER REGION, 2.2 TANDEM 3-HELIX COILED-COILS, STRUCTURAL PROTEIN				COMPLEX (DNA-BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA)	
Coumpound	HIGH MOBILITY GROUP 1 PROTEIN; CHAIN: A; DNA (5'- D(*CP*CP*(IDO) CHAIN: B; DNA (5'- CHAIN: C;	ALPHA SPECTRIN; CHAIN: A, B, C;	DNA-BINDING HIGH MOBILITY GROUP PROTEIN FRAGMENT-B (HMGB) GNA-BINDING HIME 3 HMG-BOX DOMAIN B OF RAT HMG1) (NMR, 1 STRUCTURE) 1HME 4	DNA-BINDING HIGH MOBILITY GROUP PROTEIN FRAGMENT-B (HMGB) (DNA-BINDING IHME 3 HMG-BOX DOMAIN B OF RAT HMG1) (NMR, 1 STRUCTURE) IHME 4	DNA-BNDING HIGH MOBILITY GROUP PROTEIN FRAGMENT-B (HMGB) (DNA-BINDING IHME 3 HMG-BOX DOMAIN B OF RAT HMG1) (NMR, 1 STRUCTURE) IHME 4	HUMAN SRY; IHRY 6 CHAIN: A; IHRY 7 DNA; IHRY 9 CHAIN: B; IHRY 10	HUMAN SRY; 1HRY 6 CHAIN: A; 1HRY 7 DNA; 1HRY 9 CHAIN: B; 1HRY 10	DNA-BINDING HIGH MOBILITY GROUP PROTEIN I (HMGI) BOX 2,
SeqFold								
PMF	0.01	0	0.57	0.43	_	0.39	0.78	0.21
Verify	-0.01	-0.16	0	-0.19	0.04	-0.52	0.19	0.44
PSI- BLAST	1.40E-05	7.20E-08	3.60E-14	0.00036	3.60E-13 .	9.00E-15	1.80E-16	9.00E-12
End	065	585	160	303	818	180	433	160
Start AA	530	382	107	257	369	01-	368	112
Chain	ď	∢				∢	∢	
PDB ID	lckt	lcun	o o	Ihm e	c c	1hry	1hry	lhsm
SEQ NO P	865	865	865	865	865	865	865	865

PDB Chain Start End FSI- Verify PMF SeqFold									_
PDB Chain Start End FSI- Verify PMF SeqFold	PDB annotation				GENE REGULATIONDNA HWG-D; PROTEIN-DNA COMPLEX, HMG DOMAIN, NON-SEQUENCE SPECIFIC 2 CHROMOSOMAL PROTEIN HMG-D	GENE REGULATIONDNA HMG-D; PROTEIN-DNA COMPLEX, HMG DOMAIN, NON-SEQUENCE SPECIFIC 2 CHROMOSOMAL PROTEIN HMG-D	GENE REGULATION/DNA LEF-1 HMG: LEF1, HMG, TCR-A, TRANSCRIPTION FACTOR, DNA BINDING, DNA 2 BENDING, COMPLEX (HMG DOMAIN/DNA), GENE REGULATION/DNA	GENE REGULATION/DNA LEF-1 HMG; LEF1, HMG, TCR-A, TRANSCRIPTION FACTOR, DNA BINDING, DNA 2 BENDING, COMPLEX (HMG DOMAIN/DNA), GENE REGULATION/DNA	GENE REGULATION/DNA LEF-1 HMG;
PDB Chain Start End PSJ- Verify PMF	Coumpound	COMPLEXED WITH 1HSM 3 MERCAPTOETHANOL (NMR, MINIMIZED AVERAGE STRUCTURE) 1HSM 4	DNA-BINDING HIGH MOBILITY GROUP PROTEIN (HMGI) BOX 2, COMPLEXED WITH IHSM 3 MERCAPTOETHANOL (NMR, MINIMIZED AVERAGE STRUCTURE) 14SM 4	DNA-BINDING HIGH MOBILITY GROUP PROTEIN 1 (HMG1) BOX 2, COMPLEXED WITH 1HSM 3 MERCAPTOETHANOL (NMR, MINIMIZED AVERAGE STRUCTURE) HSM 4	DNA (5:- D(*GP*CP*GP*AP*TP*AP*TP P(*GP*C)-37; CHAIN: C, D; HIGH MOBILITY GROUP PROTEN D: CHAIN: A, B;	DNA (5'- D(*GP*CP*GP*AP*TP*AP*TP *CP*GP*C)3'); CHAIN: C, D; HIGH MOBILITY GROUP PROTEIN D: CHAIN: A, B;	LYMPHOID ENHANCER- BINDING FACTOR; CHAIN: A; DNA (5'- CHAIN: B; DNA (5'- CHAIN: C;	LYMPHOID ENHANCER- BINDING FACTOR; CHAIN: A; DNA (5'- CHAIN: B; DNA (5'- CHAIN: C;	LYMPHOID ENHANCER-
PDB Chain Start End PSI- Verify	SeqFold score								
PDB Chain Start End FSI- Y	PMF score	·	0.37		0.01	0.78	0.36	0.58	1
PDB Chain Start End AA AA AA AA AA AA AA	Verify score		0.02	0.6	-0.69	0.18	-0.23	11.0	0.76
PDB Chain Start DDB D AA D D AA D D AA D D	PSI- BLAST		0.00036	5.40E-13	1.10E-10	5.40E-12	1.80E-14	3.60E-07	1.10E-14
1 hsm 1 hsm 1 hsm 1 hsm 1 dgrv A 1 dgrv A 2 lef	End		303	418	153	411	180	320	433
1 hsm 1 hsm	Start		262	369	112	367	113	262	371
	Chain				< _	A	¥ .	«	\ <u></u>
SEQ NO: NO: NO: NO: NO: NO: NO: NO: NO: NO:	PDB TD		1hsm	Ihsm	lqrv	1qrv	2lef	2lef	2lef
	SEQ ID	Ö	865	865	865	865	865	865	865

				_						_	
PDB annotation	LEFI, HMG, TCR-A, TRANSCRIPTION FACTOR, DNA BINDING, DNA 2 BENDING, COMPLEX (HMG DOMAIN/DNA), GENE REGULATION/DNA	GENE REGULATIONDNA LEF-1 HMG; LEFI, HMG, TCR-A, TRANSCRIPTION FACTOR, DNA BINDING, DNA 2 BENDING, COMPLEX (HMG DOMAINDNA), GENE REGULATION/DNA	COMPLEX (TRANSDUCER/TRANSDUCTION) GT BETA-GAMMA; MEKA, PP33; PHOSDUCIN, TRANSDUCIN, BETA- GAMMA, SIGNAL TRANSDUCTION, 2 REGULATION, PHOSPHORYLATION, G PROTEINS, THIOREDOXIN, 3 VISION, MEKA, COMPLEX (TRANSDUCER/TRANSDUCTION)		ENDOCYTOSIS/EXOCYTOSIS SYNAPTOTAGMIN ASSOCIATED 35 KDA PROTEIN, P35A, THREE HELIX BUNDLE	CONTRACTILE PROTEIN TRIPLE- HELIX COILED COIL, CONTRACTILE PROTEIN	ISOMERASE ISOMERASE, MUTASE, INTRAMOLECULAR TRANSFERASE	ISOMERASE ISOMERASE, MUTASE, INTRAMOLECULAR TRANSFERASE	TRANSCRIPTION REGULATION SIGMA70; RNA POLYMERASE SIGMA FACTOR, TRANSCRIPTION REGULATION		TRANSFERASE GLYCOSYLTRANSFERASE
Coumpound	BINDING FACTOR; CHAIN: A; DNA (5'- CHAIN: B; DNA (5'- CHAIN: C;	LYMPHOID ENHANCER- BINDING FACTOR; CHAIN: A; DNA (\$'-CHAIN: B; DNA (\$'-CHAIN: C;	TRÁNSDUCIN; CHAIN: B, G; PHOSDUCIN; CHAIN: P;	,	SYNTAXIN-IA; CHAIN: A, B, C;	HUMAN SKELETAL MUSCLE ALPHA-ACTININ 2; CHAIN: A;	METHYLMALONYL-COA MUTASE; CHAIN: A, B, C, D;	METHYLMALONYL-COA MUTASE; CHAIN: A, B, C, D;	RNA POLYMERASE PRIMARY SIGMA FACTOR; CHAIN: NULL;		SPORE COAT POLYSACCHARIDE BIOSYNTHESIS PROTEIN CHAIN: A;
SeqFold score											
PMF		89.0	-0.19		-0.2	-0.19	-0.2	-0.19	-0.19		0.22
Verify score		0.28	0.78		0.53	0.58	0.36	0.64	0.09		90.00
PSI- BLAST		3.60E-06	3.60E-12		1.80E-09	5.40E-09	3.60E-12	1.80E-10	1.80E-08		0.0018
End AA		290	709		465	467	464	482	463		302
Start AA		532	633		396	. 396	390	396	386		97
Chain		V V	<u>α</u>		₹	∢ .	٧	Ą			¥
PDB		2lef	2trc		lez3	lquu	lreq	lreq	1sig		lqgq
SEQ B S		865	865		998	998	998	998	866		871

PDB annotation	PNA-RINDING PROTEIN/RNA TRA	PRE-MENA; SPLICING REGULATION, RNP DOMAIN, RNA COMPLEX	GENE REGULATIONRNA POLY(A) BINDING PROTEIN 1, PABP 1; RRM, PROTEIN-RNA COMPLEX, GENE REGULATIONRNA	GENE REGULATION/RNA POLY(A) BINDING PROTEIN 1, PABP 1; RRM, PROTEIN-RNA COMPLEX, GENE REGULATION/RNA	GENE REGULATIONRNA POLY(A) BINDING PROTEIN 1, PABP 1; RRM, PROTEIN-RNA COMPLEX, GENE REGULATIONRNA	GENE REGULATIONRNA POLY(A) BINDING PROTEIN I, PABP I; RRM, PROTEIN-RNA COMPLEX, GENE REGULATION/RNA	RNA BINDING PROTEIN RNA- BINDING DOMAIN	NUCLEAR PROTEIN HETEROGENEOUS NUCLEAR REGONUCLEOPROTEIN A1, NUCLEAR PROTEIN, HNRNP, RBD, RRM, RNP, RNA BINDING, 2 RIBONUCLEOPROTEIN	
Coumpound	SAL LETTAL DEOTERM:	SAL-LEING, FOUTER, CALAIN, A.B.; RN4 (5"- R(P*GP*UP*UP*GP*UP*UP*U P*UP*UP*UP*UP*U)- CHAIN: P. O:	POLYDENYLATE BINDING PROTEIN 1; CHAIN: A, B, C, D, E, F, G, H; RNA (5*- R(*AP*AP*AP*AP*AP*AP*AP *AP*AP*AP*A); CHAIN: M, N, O, P, Q, K, S, T;	POLYDENYLATE BINDING PROTEIN 1; CHAIN: A, B, C, D, E, F, G, H; RNA (5'- R(*AP*AP*AP*AP*AP*AP *AP*AP*AP*A); CHAIN: M, N, O, P, Q, K, S, T;	POLYDENYLATE BINDING PROTEIN I; CHAIN: A, B, C, D, E, F, G, H; RNA (5'- R(*AP*AP*AP*AP*AP*AP*AP *AP*AP*AP*AP*3); CHAIN: M, N, O, P, Q, R, S, T;	POLYDENYLATE BINDING PROTEIN I; CHAIN: A, B, C, D, E, F, G, H; RNA (5'- R(*AP*AP*AP*AP*AP*AP *AP*AP*AP*AP*AP*AP*AP M, N, O, P, Q, R, S, T;	HU ANTIGEN C; CHAIN: A;	HNRNP A1; CHAIN: NULL;	RNA-BINDING PROTEIN SEX-LETHAL PROTEIN (C- TERMINUS, OR SECOND
SeqFold									
PMF		_	0.92	ļ		<u>-</u>	<u>-</u>		0.92
Verify	63,0	0.52	0.52	0.99	0.92	0.95	0.83	0.48	0.56
PSI- BLAST		1.40E-21	6.40E-24	1.60E-20	1.60E-20	1.60E-20	1.10E-21	3.20E-22	1.40E-21
End	3	<u>\$</u>	110	106	106	901	106	124	109
Start		o	9	27	27	27	22	21	24
Chain ID		∢	4	œ	(tr	н	V.		
PDB		167f	lcvj	lcvj	lcvj	lcvj	148z	lhai	1sxl
SEQ D NO:		872	872	872	872	872	872	872	872

PDB annotation		COMPLEX (RIBONUCLEOPROTEIN/DNA) HNRNP A1, UP1; COMPLEX (RIBONUCLEOPROTEIN/DNA), HETEROGENEOUS NUCLEAR 2 RIBONUCLEOPROTEIN A1	RNA BINDING DOMAIN RNA BINDING DOMAIN, RBD, RNA RECOGNITION MOTIF, RRM, 2 SPLICING INHIBITOR, TRANSLATIONAL INHIBITOR, SEX 3 DETERMINATION, X CHROMOSOME DOSAGE COMPENSATION	CALCIUM/PHOSPHOLIPID BINDING PROTEIN PII, CALPACTIN LIGHT CHAIN; S100 FAMILY, EF-HAND PROTEIN, LIGAND OF ANNEXIN II, 2 CALCIUM/PHOSPHOLIPID BINDING PROTEIN	CALCIUM-BINDING CNTNC; CALCIUM-BINDING, REGULATION, TROPONIN C, CARDIAC MUSCLE 2 CONTRACTION	CALCIUM BINDING CALCIUM BINDING	METAL BINDING PROTEIN CAVP; EF- HAND FAMILY, CALCIUM BINDING PROTEIN, NMR		
Coumpound	RNA-BINDING DOMAIN 1SXL 3 (RBD-2), RESIDUES 199 - 294 PLUS N-TERMINAL MET) 1SXL 4 (NMR, 17 STRUCTURES) 1SXL 5	HETEROGENEOUS NUCLEAR RIBONUCLEOPROTEIN A1; CHAIN: A12-NUCLEOTIDE SINGLE-STRANDED TELOMETRIC DNA; CHAIN: B;	SEX-LETHAL; CHAIN: A, B, C;	S100A10; CHAIN: A, B;	CARDÍAC N-TROPONIN C; CHAIN: NULL;	CALCIUM-BINDING PROTEIN; CHAIN: NULL;	CALCIUM VECTOR PROTEIN; CHAIN: A;	CALCIUM-BINDING PROTEIN CALBINDIN D9K (INTACT FORM) (NMR, 13 STRUCTURES) 1CB1 3	CALCIUM-BINDING PROTEIN CALMODULIN (VERTEBRATE) 1CLL 3
SeqFold score									+
PMF		0.99	66.0		0.29	0.15	0.16		0.16
Verify score		0.68	0.58	0.66	-0.25	-0.05	-0.22	0.35	-0.14
PSI- BLAST		1.30E-22	1.60E-20	7.20E-18	9.00E-07	5.40E-07	7.20E-07	9.00E-22	3.60E-07
End		124	104	93	7.7	78	78	8	92
Start AA		61	٥	=	21	22	22	∞	11
Chain		∢	· •	4			∢		
PDB		2up1	3xd	1а4р	lap4	1bu3	lc7w	1cb1	1cll
S B SE		872	872	873	873	873	873	873	873

PDB annotation	METAL BINDING PROTEIN YEAST FREQUENIN EF-HAND, CALCIUM	METAL TRANSPORT MRP8, S100A8, CALGRANULIN A CALCIUM-BINDING PROTEIN, CRYSTAL STRUCTURE, MAD, MIGRATION 2 INHIBITORY FACTOR. RELATED PROTEIN 8, S100 PROTEIN	COMPLEX (LIGAND/ANNEXIN) CALGIZZARIN; SIOB FAMILY, EF- HAND PROTEIN, COMPLEX (LIGAND/ANNEXIN), 2 LIGAND OF ANNEXIN II, CALCIUM/PHOSPHOLIPID BINDING PROTEIN	HYDROLASE CEREBROSIDE-3- SULFATE-SULFATASE; CEREBROSIDE-3-SULFATE HYDROLYSIS, LYSOSOMAL ENZYME, 2 HYDROLASE		LYASE DELTA3,5,DELTA2,4- DIENOYL-COENZYME A ISOMERASE, LYASE, DIENOYL-COA ISOMERASE	LYASE DELTA3,5,DELTA2,4- DIENOYL-COENZYME A ISOMERASE, LYASE, DIENOYL-COA ISOMERASE	LYASE METHYLMALONYL COA, DECARBOXYLASE	LYASE DEHALOGENASE; LYASE	LYASE CROTONASE, ENOYL-COA HYDRATASE I: LYASE, HYDRATASE, B-OXIDATION, FATTY ACID DEGRADATION, COA, 2 LIGAND BINDING
Coumpound	CALCTUM-BINDING PROTEIN NCS-1; CHAIN: A;	MIGRATION INHIBITORY FACTOR-RELATED PROTEIN 8; CHAIN: A, B;	S100C PROTEIN; CHAIN: A; ANNEXIN I; CHAIN: D;	ARYLSULFATASE A; CHAIN: NULL;		DIENOYL-COA ISOMERASE; CHAIN: A, B, C;	DIENOYL-COA ISOMERASE; CHAIN: A, B, C;	METHYLMALONYL COA DECARBOXYLASE; CHAIN: A, B, C;	4-CHLOROBENZOYL COENZYME A DEHALOGENASE; CHAIN: A, B, C;	2-ENOYL-COA HYDRATASE; CHAIN: A, B, C, D, E, F;
SeqFold										
PMF score	0.01	0.81	-	0.46		0.17	0.35	0.01	-0.11	.0.13
Verify	-0.07	0.15	0.38	0.29		-0.17	-0.02	-0.23	0.3	-0.11
PSI- BLAST	9.00E-08	1.80E-17	1.80E-18	9.60E-65		1.80E-14	3.20E-22	1.60E-18	4.80E-21	3.20E-24
End	81	83	25	330		121	44.	145	144	441
Start	=		∞	39	L	84	\$\$	54	47	47
Chain ID	A	V	.			<	∢	∢	4	4
PDB U	1fpw	1mr8	1qis	lauk		Idei	1dci	1cf8	lnzy	2dub
SEQ	873	873	873	874		875	875	875	875	875

PDB annotation	UBIQUITIN CONJUGATION UBC2; UBIQUITIN CONJUGATION, UBIQUITIN-CONJUGATING ENZYME	UBIQUITIN CONJUGATION UBC2; UBIQUITIN CONJUGATION, UBIQUITIN-CONJUGATING ENZYME	LIGASE UBIQUITIN, UBIQUITIN- CONJUGATING ENZYME, YEAST	LIGASE UBIQUITIN, UBIQUITIN- CONJUGATING ENZYME, YEAST	LIGASE UBIQUITIN, UBIQUITIN- CONJUGATING ENZYME, YEAST	SIGNALLING COMPLEX RACI; P67PHOX; SIGNALLING COMPLEX,	GTPASE, NADPH OXIDASE, PROTEIN- PROTEIN 2 COMPLEX, TPR MOTIF		CHAPERONE HOP, TPR-DOMAIN, PEPTIDE-COMPLEX, HELICAL	REPEAT, HSP90, 2 PROTEIN BINDING	CHAPERONE HOP, TPR-DOMAIN,	REPETIDE-COMPLEX, HELICAL REPEAT, HSC70, 2 HSP70, PROTEIN	BINDING SICHAI DIO DEOTETH DEDOVISAODE	RECEPTOR 1, PTS1-BP, PEROXIN-5,	PTS1 PROTEIN-PEPTIDE COMPLEX,	TETRATRICOPEPTIDE REPEAT, TPR, 2 HELICAL REPEAT	SIGNALING PROTEIN PEROXISMORE	RECEPTOR 1, PTS1-BP, PEROXIN-5,	TETT ATTICOPERTINE DEBEAT TOP 2	HELICAL REPEAT		OXIDOREDUCTASE OXIDOREDUCTASE TROPANE	ALKALOID BIOSYNTHESIS,	TROPINE, SHORT-CHAIN
Coumpound	UBIQUITIN-CONJUGATING ENZYME RAD6; CHAIN: A, B, C;	UBIQUITIN-CONJUGATING ENZYME RAD6; CHAIN: A, B, C;	UBIQUITIN CONJUGATING ENZYME; CHAIN: A;	UBIQUITIN CONJUGATING ENZYME; CHAIN: A;	UBIQUITIN CONJUGATING ENZYME; CHAIN: A;	RAS-RELATED C3 BOTULINUM TOXIN	SUBSTRATE 1; CHAIN: A;	FACTOR 2 (NCF-2) CHAIN: B;	TPR2A-DOMAIN OF HOP; CHAIN: A: HSP90-PEPTIDE	MEEVD; CHAIN: B;	TPR1-DOMAIN OF HOP;	CHAIN: A, B; HSC70- PEPTIDE; CHAIN: C, D;	DEPONISONAL TABORERIO	PEROXISOMAL TARGETING SIGNAL RECEPTOR:	CHAIN: A, B; PTSI-	CONTAINING PEPTIDE;	PEROXISOMAL TARGETING	SIGNAL 1 RECEPTOR:	CHAIN: A, B; PISI-	CHAINING PEPTIDE;		TROPINONE REDUCTASE-I;		
SeqFold score	139.32		226.34			_																		
PMF	-	1		_	1	-0.14			-0.15		-0.19		9	-0.18			-0.18					-1		
Verify score		0.84		16:0	0.71	0.15			0.31		0.02			9.0			10.0					0.35		
PSI- BLAST	3.60E-63	3.60E-63	9.00E-63	9.00E-63	4.80E-62	3.20E-09			1.60E-18		8.00E-09		7. 000 7	4.80E-14			1.40E-23					6.40E-29	-	
End	151	146	147	147	146	137			112		153			157			241				L	147		
Start	_	2	_	2	3	12			7		54						12					33		
Chain	V	4	¥	 	A	æ			<		V			∢			4					¥		
PDB	layz	layz	Idcd	Iqcq	lqcq	1.00 E+96			1clr		lelw			1tch			1fch					lael		
SEQ B B	877	877	877	877	877	878			878		878		9	878			878					881		

PDB annotation	DEHYDROGENASE	OXIDOREDUCTASE OXIDOREDUCTASE, TROPANE ALKALOID BIOSYNTHESIS, REDUCTION OF 2 TROPINONE TO TROPINE, SHORT-CHAIN DEHYDROGENASE	OXIDOREDUCTASE OXIDOREDUCTASE, DETOXIFICATION, METABOLISM, ALCOHOL 2 DEHYDROGENASE, DROSOPHILA LEBANONENSIS, SHORT-CHAIN 3 DEHYDROGENASES/REDUCTASES, TERNARY COMPLEX, NAD-3- PENTANONE 4 ADDUCT	OXIDOREDUCTASE NAD-DEPENDENT OXIDOREDUCTASE, SHORT-CHAIN ALCOHOL 2 DEHYDROGENASE, PCB DEGRADATION	OXIDOREDUCTASE SHORT-CHAIN DEHYDROGENASE, OXIDOREDUCTASE	OXIDOREDUCTASE SHORT-CHAIN DEHYDROGENASEREDUCTASE, BILE ACID CATABOLISM		OXIDOREDUCTASE SEPIAPTERIN REDUCTASE, TETRAHYDROBIOPTERIN, OXIDOREDUCTASE	OXIDOREDUCTASE NAPHTHOL REDUCTASE; OXIDOREDUCTASE	OXIDOREDUCTASE
Coumpound		TROPINONE REDUCTASE-1; CHAIN: A, B;	ALCOHOL DEHYDROGENASE; CHAIN: A, B;	CIS-BIPHENYL-2,3- DIHYDRODIOL-2,3- DEHYDROGENASE; CHAIN: NULL;	CARBONYL REDUCTASE; CHAIN: A, B, C, D;	7 ALPHA- HYDROXYSTEROID DEHYDROGENASE; CHAIN: A, B;	OXIDOREDUCTASE 3- ALPHA, 20-BETA- HYDROXYSTEROID DEHYDROGENASE (E.C.1.1.153) IHDC 3 COMPLEXED WITH CARBENOXOLONE IHDC 4	SEPIAPTERIN REDUCTASE; CHAIN: NULL;	TRIHYDROXYNAPHTHALEN E REDUCTASE; CHAIN: A, B;	TROPINONE REDUCTASE-II;
SeqFold score										
PMF		1	6.0	0.93	0.98	-	0.99	6.0		0.99
Verify score		0.36	0.14	0.32	0.1	0.68	0.34	0.47	0.38	0.49
PSI- BLAST		6.40E-29	1.40E-21	1.60E-24	4.80E-21	1.60E-34	1.10E-29	1.80E-20	4.80E-33	6.40E-29
End		147	129	147	148	147	148	129	147	147
Start		33	38	36	36	31	. 33	38	23	31
Chain		æ	<		∢ .	¥	v		¥	Ą
PDB		lael	1616	1bdb	lcyd	1fmc	Ihdo	loga	1ybv	2ae2
S B S		881	881	881	881	881	881	1881	881	881

L.	PDB ID	Chain D	Start AA	End AA	PSI- BLAST	Verify score	PMF	SeqFold score	Coumpound	PDB annotation
									CHAIN: A, B;	OXIDOREDUCTASE, TROPANE ALKALOID BIOSYNTHESIS, REDUCTION OF 2 TROPINONE TO PSEUDOTROPINE, SHORT-CHAIN DEHYDROGENASE
-										
	layz	∀	9	157	1.10E-45	0.49			UBIQUITIN-CONJUGATING ENZYME RAD6; CHAIN: A, B, C;	UBIQUITIN CONJUGATION UBC2; UBIQUITIN CONJUGATION, UBIQUITIN-CONJUGATING ENZYME
	layz	V	9	171	1.10E-45			90.32	UBIQUITIN-CONJUGATING ENZYME RAD6; CHAIN: A, B, C;	UBIQUITIN CONJUGATION UBC2: UBIQUITIN CONJUGATION, UBIQUITIN-CONJUGATING ENZYME
	164z	Q	10	157	6.40E-39	0.3			UBIQUITIN-PROTEIN LIGASE E34; CHAIN: A, B, C; UBIQUITIN CONJUGATING ENZYME E2: CHAIN: D:	LIGASE E6AP; UBCH7; BILOBAL STRUCTURE, ELONGATED SHAPE, E3 UBIQUITIN LIGASE, E2 2 UBIQUITIN CONIUGATING ENZYME
	lc4z	Q	10	166	6.40E-39			89.72	UBIQUITIN-PROTEIN LIGASE E3A; CHAIN: A, B, C; UBIQUITIN CONJUGATING ENZYME E2: CHAIN: D:	LIGASE E6AP; UBCH7; BILOBAL STRUCTURE, ELONGATED SHAPE, E3 UBIQUITIN LIGASE, E2 2 UBIQUITIN CONIUGATING ENZYME
	Iqq	Y	7	168	9.60E-49			87.99	UBIQUITIN CONJUGATING ENZYME; CHAIN: A;	LIGASE UBIQUITIN, UBIQUITIN- CONJUGATING ENZYME, YEAST
	Iqeq	V.	∞	157	9.60E-49	9.0	-		UBIQUITIN CONJUGATING ENZYMB; CHAIN: A;	LIGASE UBIQUITIN, UBIQUITIN- CONJUGATING ENZYME, YEAST
	lu9a	4	m	168	9.60E-43			74.26	UBC9; CHAIN: NULL;	UBIQUITIN-CONJUGATING ENZYME UBIQUITIN-CONJUGATING ENZYME; UBIQUITIN-CONJUGATING ENZYME, UBIQUITIN-DIRECTED 2 PROTEDLYSIS, CELL CYCLE CONTROL, LIGASE
	1u9a	v	رم	157	9.60E-43	0.17	-		UBC9; CHAIN: NULL;	UBIQUITIN-CONJUGATING ENZYME UBIQUITIN-CONJUGATING ENZYME; UBIQUITIN-CONJUGATING ENZYME, UBIQUITIN-DIRECTED 2 PROTEOLYSIS, CELL CYCLE CONTROL, LIGASE
_	2aak		5	157	6.40E-48	96.0	_		UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATION UBCI; UBIQUITIN CONJUGATION, LIGASE
	2aak		9	148	6.40E-48			93.12	UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATION UBC1; UBIQUITIN CONJUGATION, LIGASE
	2e2c			156	3.20E-44			81.53	UBIQUITIN CONJUGATING	UBIQUITIN CONJUGATION

PDB annotation	UBIQUITIN CONIUGATION, UBIQUITIN CARRIER PROTEIN, THIOESTER 2 BOND, LIGASE	UBIQUITIN CONTUGATION UBIQUITIN CONTUGATION, UBIQUITIN CARRIER PROTEIN, THIOESTER 2 BOND. LIGASE	UBIQUITIN CONIUGATION UBC7; UBIQUITIN CONIUGATION, LIGASE, YEAST	UBIQUITIN CONJUGATION UBC7; UBIQUITIN CONJUGATION, LIGASE, YEAST	UBIQUITIN CONJUGATION UBC2; UBIQUITIN CONJUGATION, UBIQUITIN-CONJUGATING ENZYME	UBIQUITIN CONJUGATION UBC2; UBIQUITIN CONJUGATION, UBIQUITIN-CONJUGATING ENZYME	LIGASE E6AP, UBCH7; BILOBAL STRUCTURE, ELONGATED SHAPE, E3 UBIQUITIN LIGASE, E2 2 UBIQUITIN CONJUGATING ENZYME	LIGASE E6AP, UBCH7; BILOBAL STRUCTURE, ELONGATED SHAPE, E3 UBIQUITIN LIGASE, E2 2 UBIQUITIN CONTUGATING ENZYME	LIGASE UBIQUITIN, UBIQUITIN- CONJUGATING ENZYME, YEAST	LIGASE UBIQUITIN, UBIQUITIN- CONJUGATING ENZYME, YEAST	UBIQUITIN-CONJUGATING ENZYME UBIQUITIN-CONJUGATING ENZYME, UBIQUITIN-CONJUGATING ENZYME, UBIQUITIN-DIRECTED 2 PROTEOLYSIS, CELL CYCLE CONTROL, LIGASE	UBIQUITIN-CONJUGATING ENZYME, UBIQUITIN-CONJUGATING ENZYME, UBIQUITIN-CONJUGATING ENZYME, UBIQUITIN-DIRECTED 2
Id	UBIQUITIN COUBIQUITIN COURTIN COURT THIOESTER 2	UBIQUITIN CO UBIQUITIN CO UBIQUITIN CO	UBIQUITIN C UBIQUITIN C YEAST	UBIQUITIN C UBIQUITIN C YEAST	UBIQUITIN C UBIQUITIN C	UBIQUITIN C UBIQUITIN C UBIQUITIN-C	LIGASE E6AP; UBCH7; BII STRUCTURE, ELONGATE UBIQUITIN LIGASE, E2 2 CONTUGATING ENZYME	LIGASE E6AF STRUCTURE, UBIQUITIN L	LIGASE UBIC	LIGASE UBIC	UBIQUITIN-CONUGATII UBIQUITIN-CONUGATII UBIQUITIN-CONUGATII UBIQUITIN-DIRECTED 2 PROTEOLYSIS, CELL CY CONTROL, LIGASE	UBIQUITIN-CONJUGATII UBIQUITIN-CONJUGATII UBIQUITIN-CONJUGATII UBIQUITIN-DIRECTED 2
Coumpound	ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONIUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN-CONJUGATING ENZYME RAD6; CHAIN: A, B, C;	UBIQUITIN-CONJUGATING ENZYME RAD6; CHAIN: A, B, C;	UBIQUITIN-PROTEIN LIGASE E3A; CHAIN: A, B, C; UBIQUITIN CONJUGATING ENZYME E2; CHAIN: D;	UBIQUITIN-PROTEIN LIGASE E3A; CHAIN: A, B, C; UBIQUITIN CONJUGATING ENZYME E2; CHAIN: D;	UBIQUITIN CONJUGATING ENZYME; CHAIN: A;	UBIQUITIN CONJUGATING ENZYME; CHAIN: A;	UBC9; CHAIN: NULL;	UBC9; CHAIN: NULL;
SeqFold score				121.4	114.6			104.56	104.4		94.76	
PMF		_	-			-	1			-		-
Verify score		0.27	0.63			0.73	0.4			0.53		0.8
PSI- BLAST		3.20E-44	3.20E-51	3.20E-51	6.40E-50	6.40E-50	3.20E-42	3.20E-42	1.10E-53	1.10E-53	4.80E-45	4.80E-45
End AA		157	157	165	178	178	165	169	166	166	174	173
Start AA		3	9	7	9	٥	01	10	7	∞	m	8
Chain					V	4	Ω	Ω	¥.	∢	4	<
PDB		2e2c	2ucz	2ucz	18yz	layz	1042	104z	Iqeq	1qcq	1u9a	1u9a
SEQ EQ	Ž	882	882	882	882	882	882	882	882	882	882	882

PDB annotation	PROTEOLYSIS, CELL CYCLE CONTROL, LIGASE	UBIQUITIN CONTUGATION UBCI; UBIQUITIN CONTUGATION, LIGASE	UBIQUITIN CONJUGATION UBC1; UBIQUITIN CONJUGATION, LIGASE	UBIQUITIN CONJUGATION	UBIQUITIN CARRIER PROTEIN,	THIOESTER 2 BOND, LIGASE	UBIQUITIN CONJUGATION	UBIQUITIN CONJUGATION,	THIOESTER 2 BOND, LIGASE	UBIQUITIN CONJUGATION UBC7:	UBIQUITIN CONJUGATION, LIGASE, YEAST	UBIQUITIN CONJUGATION UBC7; UBIQUITIN CONJUGATION, LIGASE,	YEAST	UBIQUITIN CONJUGATION UBC2;	UBIQUITIN CONJUGATION,	OBICOLIN-CONCOUNT TO DISCUSSION OF THE CONTOUR PROPERTY OF THE CONTOUR PROPERT	LIGASE E6AP; UBCH7; BILUBAL	SIRUCIUKE, ELONGAIED SHAFE, E3	UBIQUITIN LIGASE, EZ 2 UBIQUITIN	LIGASE E6AP; UBCH7; BILOBAL	STRUCTURE, ELONGATED SHAPE, E3	UBIQUITIN LIGASE, E2 2 UBIQUITIN	CONJUGATING ENZYME	LIGASE UBIQUITIN, UBIQUITIN- CONJUGATING ENZYME, YEAST	UBIQUITIN-CONJUGATING ENZYME	UBIQUITIN-CONJUGATING ENZYME;	UBIQUITIN-CONJUGATING ENZYME,	DESCRIPTION OF THE COLUMN TE	CONTROL, LIGASE	UBIQUITIN CONJUGATION UBCI; UBIQUITIN CONJUGATION, LIGASE
Coumpound		UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATING	ENZYME; CHAIN: NULL;		UBIQUITIN CONJUGATING	ENZYME: CHAIN: NULL;		UBIQUITIN CONJUGATING	ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;		UBIQUITIN-CONJUGATING	ENZYME RAD6; CHAIN: A, B,	اد	UBIQUITIN-PROTEIN	LIGASE E3A; CHAIN: A, B, C;	UBIQUITIN CONJUGATING FNZVMF F2: CHAIN: D:	UBIOUITIN-PROTEIN	LIGASE E3A; CHAIN: A, B, C;	UBIQUITIN CONJUGATING	ENZYME E2; CHAIN: D;	UBIQUITIN CONJUGATING ENZYME: CHAIN: A:	UBC9; CHAIN: NULL;					UBIQUITIN CONJUGATING ENZYME: CHAIN: NULL;
SeqFold score			116.78	109.63								148.79								51.09										
PMF		-					-			-				0.75			0.54							9.0	0.72					0.55
Verify score		0.62					0.55			0.59				0.19			-0.22							0.2	-0.18					90:0
PSI- BLAST		8.00E-52	8.00E-52	1.60E-45			1,60E-45			1.60E-55		1.60E-55		4.80E-37			1.60E-28			1.60E-28				3.20E-39	3.20E-34					3.20E-38
End		172	174	176			160			160		174		128			128			163				128	128					128
Start		8	و	_			3			9		7		9			<u> </u>			2				8	5					2
Chain														¥			Ω							Ą	4					
PDB		2aak	2aak	2e2c			2e2c			2ncz		2ucz		layz			1c4z			1047				lqcq	1u9a					2aak
SEQ P		882	882	882			882			882		882		882			882			882				882	882					882

					_,			_			_										_	
PDB annotation	UBIQUITIN CONJUGATION UBCI; UBIQUITIN CONJUGATION, LIGASE	UBIQUITIN CONJUGATION UBIQUITIN CONJUGATION, UBIQUITIN CARRIER PROTEIN, THIOESTER 2 BOND, LIGASE	UBIQUITIN CONJUGATION UBC7; UBIQUITIN CONJUGATION, LIGASE, YEAST	UBIQUITIN CONJUGATION UBC7; UBIQUITIN CONJUGATION, LIGASE, YEAST		UBIQUITIN CONJUGATION UBC2; UBIQUITIN CONJUGATION, UBIQUITIN-CONJUGATING ENZYME	UBIQUITIN CONJUGATION UBC2; UBIQUITIN CONJUGATION, UBIQUITIN-CONJUGATING ENZYME	LIGASE EKAP, LIBCH7: BILOBAL	STRUCTURE, ELONGATED SHAPE, E3	UBIQUITIN LIGASE, E2 2 UBIQUITIN CONJUGATING ENZYME	LIGASE E6AP; UBCH7; BILOBAL	STRUCTURE, ELONGATED SHAPE, E3 UBIOUITIN LIGASE, E2 2 UBIOUITIN	CONJUGATING ENZYME	LIGASE UBIQUITIN, UBIQUITIN- CONIIIGATING ENZYME, YEAST	LIGASE UBIQUITIN, UBIQUITIN- CONTIGATING ENZYME, YEAST	UBIQUITIN-CONJUGATING ENZYME	UBIQUITIN-CONJUGATING ENZYME;	UBIQUITIN-CONJUGATING ENZYME,	PROTEOLYSIS, CELL CYCLE	CONTROL, LIGASE	UBIQUITIN-CONJUGATING ENZYME	UBIQUITIN-CONJUGATING ENZYME; UBIQUITIN-CONJUGATING ENZYME, UBIQUITIN-DIRECTED 2
Coumpound	UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONIUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;		UBIQUITIN-CONJUGATING ENZYME RAD6; CHAIN: A, B, C;	UBIQUITIN-CONJUGATING ENZYME RAD6; CHAIN: A, B, C;	TIBIOITHIN-PROTEIN	LIGASE E3A; CHAIN: A, B, C;	UBIQUITIN CONJUGATING ENZYME E2; CHAIN: D;	UBIQUITIN-PROTEIN	LIGASE E3A; CHAIN: A, B, C; UBIOUITIN CONJUGATING	ENZYME E2; CHAIN: D;	UBIQUITIN CONJUGATING FNZYME: CHAIN: A:	UBIQUITIN CONJUGATING ENZYME: CHAIN: A:	UBC9; CHAIN: NULL;					UBC9; CHAIN: NULL:	
SeqFold score	50.62			56.09			90.32		_		89.72			87.99		74.26						
PMF score		0.25	0.49			<u> </u>		-	-						-							
Verify score		0.17	-0.09			0.49		0 3	3				_		9.0						0.17	
PSI- BLAST	3.20E-38	1.10E-35	1.60E-35	1.60E-35		1.10E-45	1.10E-45	6 40E-30	0.405-39		6.40E-39			9.60E-49	9.60E-49	9.60E-43					9.60E-43	
End	155	126	128	136		157	171	157	Ì		.166			168	157	168					157	
Start	9	м	9	7		9	٥	5	3		2			7	∞	3				_	2	
Chain ID						⋖	<		J .		Δ			∢	4	V					∢	
PDB ID	2aak	2e2c	2ucz	2ncz		layz	layz	10.43	75		1042			Jdcd	Iqcq	1u9a					lu9a	
SEQ NO.	882	882	882	882		883	883	003	ê		883			883	883	883	_				883	

PDB annotation	PROTEOLYSIS, CELL CYCLE CONTROL, LIGASE	UBIQUITIN CONJUGATION UBCI; UBIQUITIN CONJUGATION, LIGASE	UBIQUITIN CONJUGATION UBCI; UBIQUITIN CONJUGATION, LIGASE	UBIQUITIN CONJUGATION UBIQUITIN CONJUGATION, UBIQUITIN CARRIER PROTEIN, THIOESTER 2 BOND, LIGASE	UBIQUITIN CONJUGATION UBIQUITIN CONJUGATION, UBIQUITIN CARRIER PROTEIN, THIOESTER 2 BOND, LIGASE	UBIQUITIN CONJUGATION UBC7; UBIQUITIN CONJUGATION, LIGASE, YEAST	UBIQUITIN CONJUGATION UBC7; UBIQUITIN CONJUGATION, LIGASE, YEAST	UBIQUITIN CONJUGATION UBC2; UBIQUITIN CONJUGATION, UBIQUITIN-CONJUGATING ENZYME	UBIQUITIN CONJUGATION UBC2; UBIQUITIN CONJUGATION, UBIQUITIN-CONJUGATING ENZYME	LIGASE E6AP; UBCH7; BILOBAL STRUCTURE, ELONGATED SHAPE, E3 UBIQUITIN LIGASE, E2 2 UBIQUITIN CONJUGATING ENZYME	LIGASE E6AP; UBCH7; BILOBAL STRUCTURE, ELONGATED SHAPE, E3 UBIQUITIN LIGASE, E2 2 UBIQUITIN CONJUGATING ENZYME	LIGASE UBIQUITIN, UBIQUITIN- CONJUGATING ENZYME, YEAST	LIGASE UBIQUITIN, UBIQUITIN- CONJUGATING ENZYME, YEAST	UBIQUITIN-CONJUGATING ENZYME, UBIQUITIN-CONJUGATING ENZYME, UBIQUITIN-CONJUGATING ENZYME,
Coumpound		UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN-CONJUGATING ENZYME RAD6; CHAIN: A, B, C;	UBIQUITIN-CONIUGATING ENZYME RAD6; CHAIN: A, B, C;	UBIQUITIN-PROTEIN LIGASE E3A; CHAIN: A, B, C; UBIQUITIN CONJUGATING ENZYME E2; CHAIN: D;	UBIQUITIN-PROTEIN LIGASE E3A; CHAIN: A, B, C; UBIQUITIN CONJUGATING ENZYME E2; CHAIN: D;	UBIQUITIN CONJUGATING ENZYME; CHAIN: A;	UBIQUITIN CONJUGATING ENZYME; CHAIN: A;	UBC9; CHAIN: NULL;
SeqFold score			93.12	81.53			121.4	114.6			104.56	104.4		94.76
PMF		_			-	-			_	-			-	
Verify score		0.38			0.27	0.63			0.73	0.4			0.53	
PSI- BLAST		6.40E-48	6.40E-48	3.20E-44	3.20E-44	3.20E-51	3.20E-51	6.40E-50	6.40E-50	3.20E-42	3.20E-42	1.10E-53	1.10E-53	4.80E-45
End		157	148	156	157	157	165	178	178	165	691	166	166	174
Start AA		2	9	_	m	9	7	9	٥	01	01	7	∞	т ·
Chain TD								V	<	Q	Q	¥	¥	∢
PDB ID		2aak	2aak	2e2c	2e2c	2ucz	2ucz	layz	layz	104z	1c4z	1qcq	1qcq	lu9a
SEQ ID		883	883	883	883	883	883	883	883	883	883	883	883	883

PDB annotation	UBIQUITIN-DIRECTED 2 PROTEOLYSIS, CELL CYCLE CONTROL, LIGASE											CONJUGATING ENZYME, YEAST
Coumpound		UBC9; CHAIN: NULL;	UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONIUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN-CONJUGATING ENZYME RAD6; CHAIN: A, B, C;	UBIQUITIN-PROTEIN LIGASE E34; CHAIN: A, B, C; UBIQUITIN CONJUGATING ENZYME E2; CHAIN: D;	UBIQUITIN-PROTEIN LIGASE E34; CHAIN: A, B, C; UBIQUITIN CONJUGATING ENZYME E2; CHAIN: D;	UBIQUITIN CONJUGATING ENZYME; CHAIN: A;
SeqFold score				116.78	109.63			148.79			51.09	
PMF		-	1			-			0.75	0.54		9.0
Verify score		0.8	0.62			0.55	0.59		0.19	-0.22		0.2
PSI- BLAST		4.80E-45	8.00E-52	8.00E-52	1.60E-45	1.60E-45	1.60E-55	1.60E-55	4.80E-37	1.60E-28	1.60E-28	3.20E-39
End		173	172	174	176	160	160	174	128	128	163	128
Start		5	8	9	-	E .	9	7	9	01	10	œ
Chain ID		<							∢	ρ	Ω	∢
PDB		1u9a	2aak	2aak	2e2c	2e2c	2ucz	2ucz	1ayz	104z	1c4z	1400
SEQ	Ë	883	883	883	883	883	883	883	883	883	883	883

PDB annotation	UBIQUITIN-CONIUGATING ENZYME; UBIQUITIN-CONIUGATING ENZYME, UBIQUITIN-DIRECTED 2 PROTEOLYSIS, CELL CYCLE CONTROL, LIGASE	UBIQUITIN CONJUGATION UBCI; UBIQUITIN CONJUGATION, LIGASE	UBIQUITIN CONJUGATION UBCI; UBIQUITIN CONJUGATION, LIGASE	UBIQUITIN CONIUGATION UBIQUITIN CONIUGATION, UBIQUITIN CARRIER PROTEIN, THIOESTER 2 BOND, LIGASE	UBIQUITIN CONJUGATION UBC7; UBIQUITIN CONJUGATION, LIGASE, YEAST	UBIQUITIN CONJUGATION UBC7; UBIQUITIN CONJUGATION, LIGASE, YEAST	UBIQUITIN CONJUGATION UBC2; UBIQUITIN CONJUGATION, UBIQUITIN-CONJUGATING ENZYME	UBIQUITIN CONJUGATION UBC2; UBIQUITIN CONJUGATION, UBIQUITIN-CONJUGATING ENZYME	LIGASE E6AP, UBCH7, BILOBAL STRUCTURE, ELONGATED SHAPE, E3 UBIQUITIN LIGASE, E2 2 UBIQUITIN CONJUGATING ENZYME	LIGASE EGAP, UBCH7; BILOBAL STRUCTURE, ELONGATED SHAPE, E3 UBIQUITIN LIGASE, E2 2 UBIQUITIN CONJUGATING ENZYME	LIGASE UBIQUITIN, UBIQUITIN- CONJUGATING ENZYME, YEAST	LIGASE UBIQUITIN, UBIQUITIN- CONJUGATING ENZYME, YEAST	UBIQUITIN-CONIUGATING ENZYME UBIQUITIN-CONIUGATING ENZYME; UBIQUITIN-CONIUGATING ENZYME,
Coumpound		UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN-CONJUGATING ENZYME RAD6: CHAIN: A. B, C;	UBIQUITIN-CONJUGATING ENZYME RAD6; CHAIN: A, B, C;	UBIQUITIN-PROTEIN LIGASE E3A; CHAIN: A, B, C; UBIQUITIN CONJUGATING ENZYME E2; CHAIN: D;	UBIQUITIN-PROTEIN LIGASE E34; CHAIN: A, B, C; UBIQUITIN CONJUGATING ENZYME E2; CHAIN: D;	UBIQUITIN CONJUGATING ENZYME; CHAIN: A;	UBIQUITIN CONJUGATING ENZYME; CHAIN: A;	UBC9; CHAIN: NULL;
SeqFold score			50.62			56.09		90.32		89.72	87.99		74.26
PMF		0.55		0.25	0.49		-		-				
Verify score		90.0		0.17	-0.09		0.49		0.3			9.0	
PSI- BLAST		3.20E-38	3.20E-38	1.10E-35	1.60E-35	1.60E-35	1.10E-45	1.10E-45	6.40E-39	6.40E-39	9.60E-49	9.60E-49	9.60E-43
End		128	155	126	128	136	157	171	157	166	168	157	168
Start		5	9	3	9	7	9	9	01 ·	10	7	∞	£
Chain							4	∢	Q	Δ	¥	V	V
PDB ID		2aak	2aak	2 e 2c	2ucz	2ucz	layz	layz	1c4z	1c4z	1qcq	Iqeq	lu9a
SEQ NO		883	883	883	883	883	884	884	884	884	884	884	884

PDB annotation	UBIQUITIN-DIRECTED 2 PROTEOLYSIS, CELL CYCLE CONTROL, LIGASE	UBIQUITIN-CONJUGATING ENZYME UBIQUITIN-CONUGATING ENZYME; UBIQUITIN-CONJUGATING ENZYME, UBIQUITIN-DIRECTED 2 PROTEOLYSIS, CELL CYCLE CONTROL, LIGASE	UBIQUITIN CONJUGATION UBC1; UBIQUITIN CONJUGATION, LIGASE	UBIQUITIN CONJUGATION UBCI; UBIQUITIN CONJUGATION, LIGASE	UBIQUITIN CONJUGATION UBIQUITIN CONJUGATION, UBIQUITIN CARRIER PROTEIN, THIOESTER 2 BOND, LIGASE	UBIQUITIN CONIUGATION UBIQUITIN CONIUGATION, UBIQUITIN CARRIER PROTEIN, THIOESTER 2 BOND, LIGASE	UBIQUITIN CONTUGATION UBC7; UBIQUITIN CONTUGATION, LIGASE, YEAST	UBIQUITIN CONJUGATION UBC7; UBIQUITIN CONJUGATION, LIGASE, YEAST	UBIQUITIN CONJUGATION UBC2; UBIQUITIN CONJUGATION, UBIQUITIN-CONJUGATING ENZYME	UBIQUITIN CONJUGATION UBC2; UBIQUITIN CONJUGATION, UBIQUITIN-CONJUGATING ENZYME	LIGÁSE E6AP; UBCH?; BILOBAL STRUCTURE, ELONGATED SHAPE, E3 UBIQUITIN LIGASE, E2 2 UBIQUITIN CONJUGATING ENZYME	LIGASE E6AP; UBCH7; BILOBAL STRUCTURE, ELONGATED SHAPE, E3 UBIQUITIN LIGASE, E2 2 UBIQUITIN CONIUGATING ENZYME
Coumpound		UBC9; CHAIN: NULL;	UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONIUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN-CONJUGATING ENZYME RAD6: CHAIN: A, B. C;	UBIQUITIN-CONIUGATING ENZYME RAD6; CHAIN: A, B, C;	UBIQUITIN-PROTEIN LIGASE E3A; CHAIN: A, B, C; UBIQUITIN CONJUGATING ENZYME E2; CHAIN: D;	UBIQUITIN-PROTEIN LIGASE E3A; CHAIN: A, B, C; UBIQUITIN CONJUGATING ENZYME E2; CHAIN: D;
SeqFold.				93.12	81.53			121.4	114.6			104.56
PMF score		_	-			_	_			-	-	
Verify score		0.17	0.38			0.27	0.63			0.73	0.4	
PSI. BLAST		9.60E-43	6.40E-48	6.40E-48	3.20E-44	3.20E-44	3.20E-51	3.20E-51	6.40E-50	6.40E-50	3.20E-42	3.20E-42
End AA		157	157	148	156	157	157	165	178	178	165	691
Start AA		8	5	9	-	E	9	7	9	9	9	10
Chain ID		V							∢	∢	Q	Q
PDB DD		lu9a	2aak	2aak	2620	2 e 2c	2ucz	2ucz	layz	layz	1c4z	1c4z
SEQ EQ		884	884	884	884	884	884	884	884	884	884	884

							_				_	_			_		_		r-			_		_				1		\neg
PDB annotation	LIGASE UBIQUITIN, UBIQUITIN- CONJUGATING ENZYME, YEAST	LIGASE UBIQUITIN, UBIQUITIN- CONJUGATING ENZYME, YEAST	UBIQUITIN-CONJUGATING ENZYME	UBIQUITIN-CONJUGATING ENZYME;	TRICOLLING CONTROL ENGINE,	UBIQUITIN-DINECTED 2 DECTEO! VOIC OF! 1 CVC! F	CONTROL, LIGASE	UBIQUITIN-CONJUGATING ENZYME	UBIQUITIN-CONJUGATING ENZYME;	UBIQUITIN-CONJUGATING ENZYME,	PROTEOLYSIS, CELL CYCLE	CONTROL, LIGASE	UBIQUITIN CONJUGATION UBCI; UBIOUITIN CONJUGATION, LIGASE	UBIQUITIN CONJUGATION UBCI;		TEICULIN CONTIGATION	TIBIOLITIN CARRIER PROTEIN	THIOESTER 2 BOND, LIGASE	UBIOUTTIN CONJUGATION	UBIQUITIN CONJUGATION,	UBIQUITIN CARRIER PROTEIN,	THIOESTER 2 BOND, LIGASE	UBIQUITIN CONJUGATION UBC7; UBIQUITIN CONJUGATION, LIGASE,	YEASI	UBIQUITIN CONJUGATION UBC7; UBIQUITIN CONJUGATION, LIGASE, YEAST	UBIQUITIN CONJUGATION UBC2;	UBIQUITIN CONJUGATION,	LIGASE FEAP URCH7: BILOBAL	STRUCTURE, ELONGATED SHAPE, E3	CONJUGATING ENZYME
Coumpound	UBIQUITIN CONJUGATING ENZYME; CHAIN: A;	UBIQUITIN CONJUGATING ENZYME; CHAIN: A;	UBC9; CHAIN: NULL;					UBC9; CHAIN: NULL;					UBIQUITIN CONJUGATING ENZYME: CHAIN: NULL:	UBIQUITIN CONJUGATING	ENZIME, CHAIN: NOLL,	UBIQUITIN CONJUGATING	ENGINE, CIRCIN, NOED,		UBIOUITIN CONJUGATING	ENZYME: CHAIN: NULL:			UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;		UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN-CONJUGATING	ENZYME RAD6; CHAIN: A, B,	TIBIOTITH PROTEIN	LIGASE E3A; CHAIN: A, B, C;	UBIQUITIN CONJUGATING ENZYME E2; CHAIN: D;
SeqFold score	104.4		94.76											116.78	,	109.63									148.79		-			
PMF		1						_											-				-			0.75		2	<u> </u>	
Verify score		0.53						8.0					0.62	_					0 55	}			0.59			0.19		0,00	7	
PSI-	1.10E-53	1.10E-53	4.80E-45					4.80E-45					8.00E-52	8.00E-52		1.60E-45			1 60F-45	200:1			1.60E-55		1.60E-55	4.80E-37		1 605 38	1.005-20	
End	991	166	174					173					172	174		176			160	3			160		174	128		120	971	
Start AA	7	∞	3					5					2	9		<u>-</u> _			-	•			9		7	9		2	2	
Chain	A	4	V					\		-																<			2	
PDB DD	Iqcd	Iqq	1198		_			lu9a					2aak	2aak		2e2c			2000	777	_		2ncz		2ncz	layz		100	75.2	
SEQ D C	884	884	884					884	}				884	884		884			884	5			884		884	884		700	60	

	E Z		6 6, 6,	ய	ய		ជា	ng	1			пí	-
PDB annotation	LIGASE E6AP, UBCH7, BILOBAL STRUCTURE, ELONGATED SHAPE, E3 UBIQUITIN LIGASE, E2 2 UBIQUITIN CONJUGATING ENZYME	LIGASE UBIQUITIN, UBIQUITIN- CONJUGATING ENZYME, YEAST	UBIQUITIN-CONJUGATING ENZYME UBIQUITIN-CONJUGATING ENZYME; UBIQUITIN-DIRECTED 2 UBIQUITIN-DIRECTED 2 PROTEOL YSIS, CELL CYCLE CONTROL, LIGASE	UBIQUITIN CONJUGATION UBCI; UBIQUITIN CONJUGATION, LIGASE	UBIQUITIN CONJUGATION UBCI; UBIQUITIN CONJUGATION, LIGASE	UBIQUITIN CONJUGATION UBIQUITIN CONJUGATION, UBIQUITIN CARRIER PROTEIN, THIOESTER 2 BOND, LIGASE	UBIQUITIN CONIUGATION UBC7; UBIQUITIN CONIUGATION, LIGASE, YEAST	UBIQUITIN CONIUGATION UBC7; UBIQUITIN CONIUGATION, LIGASE, YEAST	LIPID TRANSPORT APO A-I: LIPOPROTEIN, LIPID TRANSPORT, CHOLESTEROL METABOLISM, 2 ATHEROSCLEROSIS, HDI, LCAT- ACTIVATION	HYDROLASE ERA, GTPASE, RNA- BINDING, RAS-LIKE, HYDROLASE	TRANSLATION TRANSLATIONAL GTPASE	CONTRACTILE PROTEIN TRIPLE. HELIX COILED COIL, CONTRACTILE PROTEIN	
	LIGASE E6A STRUCTUR UBIQUITIN CONJUGATI	LIGASE UBI CONTUGATI	UBIQUITIN-CONJUC UBIQUITIN-CONJUC UBIQUITIN-DIRECT PROTEOLYSIS, CEL CONTROL, LIGASE	UBIQUITIN UBIQUITIN	VIITINOIBIN VIITINOIBIN	UBIQUITIN UBIQUITIN UBIQUITIN THIOESTER	UBIQUITIN UBIQUITIN YEAST	UBIQUITIN UBIQUITIN YEAST	LIPID TRANSI LIPOPROTEIN CHOLESTERC ATHEROSCLE	HYDROLAS BINDING R	TRANSLATI GTPASE	CONTRACT HELIX COIL	
Coumpound	UBIQUITIN-PROTEIN LIGASE E3A; CHAIN: A, B, C; UBIQUITIN CONJUGATING ENZYME E2; CHAIN: D;	UBIQUITIN CONJUGATING ENZYME; CHAIN: A;	UBC9; CHAIN: NULL;	UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONIUGATING ENZYME; CHAIN: NULL;	APOLIPOPROTEIN A-1; CHAIN: A, B, C, D;	GTP-BINDING PROTEIN ERA; CHAIN: A. B:	TRANSLATION INITIATION PACTOR IP2/EIF5B; CHAIN: A:	HUMAN SKELETAL MUSCLE ALPHA-ACTININ 2; CHAIN:	
SeqFold score	51.09				50.62			56.09	53.53				
PMF score		9.6	0.72	0.55		0.25	0.49			0.01	0.03	-0.01	
Verify score		0.2	-0.18	90.0		0.17	-0.09			-0.14	-0.34	0.23	
PSI- BLAST	1.60E-28	3.20E-39	3.20E-34	3.20E-38	3.20E-38	1.10E-35	1.60E-35	1.60E-35	5.40E-07	1.60E-37	6.40E-06	1.80E-08	
End	163	128	128	128	155	126	128	136	427	350	289	426	•
Start AA		∞	w	s	9	<u>8</u>	و	7	237	39	45	283	
Chain D	Д	<	∢						<	<	¥	¥	
PDB ID	lc4z	1969	1u9a	2aak	2aak	2e2c	2ucz	2ucz	lavI	lega	1g7s	lquu	-
SEQ ID No:	884	884	. 884	884	884	884	884	884	885	885	885	882	

PDB annotation	HELIX COILED COIL, CONTRACTILE PROTEIN	LIPID TRANSPORT APO A-1; LIPOPROTEIN, LIPID TRANSPORT, CHOLESTEROL METABOLISM, 2 ATHEROSCLEROSIS, HDL, LCAT- ACTIVATION	HYDROLASE ERA, GTPASE, RNA- BINDING, RAS-LIKE, HYDROLASE	TRANSLATION TRANSLATIONAL GTPASE	CONTRACTILE PROTEIN TRIPLE- HELIX COILED COIL, CONTRACTILE PROTEIN	CONTRACTILE PROTEIN TRIPLE- HELIX COILED COIL, CONTRACTILE PROTEIN	IMMUNOGLOBULIN IMMUNOGLOBULIN, KAPPA LIGHT- CHAIN DIMER HEADER	COMPLEX (ANTIBODY/ANTIGEN) FAB-12; VEGF; COMPLEX (ANTIBODY/ANTIGEN), ANGIOGENIC FACTOR	COMPLEX (HUMANIZED ANTIBODY/HYDROLASE) MURAMIDASE; HUMANIZED ANTIBODY, ANTIBODY COMPLEX, FV, ANTI-LYSOZYME, 2 COMPLEX (HUMANIZED ANTIBODY/HYDROLASE)	IMMUNE SYSTEM REIV, STABILIZED IMMUNOGLOBULIN FRAGMENT, BENCE-JONES 2 PROTEIN, IMMUNE SYSTEM	IMMUNE SYSTEM FAB-IBP COMPLEX CRYSTAL STRUCTURE 2.7A RESOLUTION BINDING 2 OUTSIDE
Coumpound	ALPHA-ACTININ 2; CHAIN: A;	APOLIPOPROTEIN A-1; CHAIN: A, B, C, D;	GTP-BINDING PROTEIN ERA; CHAIN: A, B;	TRANSLATION INITIATION FACTOR IF2/EIF5B; CHAIN: A;	HUMAN SKELETAL MUSCLE ALPHA-ACTININ 2; CHAIN: A:	HUMAN SKELETAL MUSCLE ALPHA-ACTININ 2; CHAIN: A;	IMMUNOGLOBULIN; CHAIN: A, B;	FAB FRAGMENT; CHAIN: L, H, J, K; VASCULAR ENDOTHELIAL GROWTH FACTOR; CHAIN: V, W;	HULYS11; CHAIN: A, B, D, E: LYSOZYME; CHAIN: C, F;	IG KAPPA CHAIN V-I REGION REI; CHAIN: A, B;	IGM RF 2A2; CHAIN: A, C, E; IGM RF 2A2; CHAIN: B, D, F; IMMUNOGLOBULIN G
SeqFold score		53.53									
PMF			10.0	0.03	-0.01	-0.01	0.17	0.11	-0.09	0.34	.0.1
Verify			-0.14	-0.34	0.23	0.23	0.2	0.13	0.18	0.15	-0.2
PSI- BLAST		5.40E-07	1.60E-37	6.40E-06	1.80E-08	1.80E-08	3.20B-41	3.20E-43	1.60E-42	4.80E-43	3.20E-43
End		427	350	289	426	445	118	116	119	118	115
Start		237	39	45	283	302	61	19	19	61	61
Chain		4	<	\	<	∀	¥	i)	∢	<	∢
PDB ID		lavl	lega	1g7s	Iquu	lquu	1b6d	ıbjı	1bvk	N W	Idee
SEQ S B S		988	988	886	886	886	887	887	887	887	887

					, 	_					
PDB annotation	THE ANTIGEN COMBINING SITE SUPERANTIGEN FAB VH3 3 SPECIFICITY			IMMUNOGLOBULIN TR1.9, ANTI- THYROID PEROXIDASE, AUTOANTIBODY, 2 IMMUNOGLOBULIN			DNA-BINDING PROTEIN ISL-1HD DNA-BINDING PROTEIN, HOMEODOMAIN, LIM DOMAIN	COMPLEX (DNA-BINDING PROTEIN/DNA) DNA-BINDING PROTEIN, DNA, PAIRED BOX, TRANSCRIPTION 2 REGULATION	COMPLEX (DNA-BINDING PROTEIN/DNA) DNA-BINDING PROTEIN, DNA, PAIRED BOX, TRANSCRIPTION 2 REGULATION	COMPLEX (DNA-BINDING PROTEIN/DNA) DNA-BINDING PROTEIN, DNA, PAIRED BOX, TRANSCRIPTION 2 REGULATION	COMPLEX (DNA-BINDING PROTEIN/DNA) DNA-BINDING PROTEIN, DNA, PAIRED BOX,
Coumpound	BINDING PROTEIN A; CHAIN: G, H:	IMMUNOGLOBULIN FV FRAGMENT OF A HUMANIZED VERSION OF HE ANTI-CD18 1FGV 3 ANTIBODY 'H52' (HUH52-AA FV) 1FGV 4	IMMUNOGLOBULIN IMMUNOGLOBULIN M (IG- M) FV FRAGMENT 11GM 3	TRI.9 FAB; CHAIN: L, H;	IMMUNOGLOBULIN FAB FRAGMENT OF A HUMANIZED VERSION OF THE ANTI-CD18 2FGW 3 ANTIBODY 'H52' (HUH52-OZ FAB) 2FGW 4		INSULIN GENE ENHANCER PROTEIN ISL-1; CHAIN: NULL;	PAIRED PROTEIN; CHAIN: A, B, C; DNA; CHAIN: D, E, F	PAIRED PROTEIN; CHAIN: A, B. C; DNA; CHAIN: D, E, F	PAIRED PROTEIN; CHAIN: A, B, C; DNA; CHAIN: D, E, F	PAIRED PROTEIN; CHAIN: A, B, C; DNA; CHAIN: D, E, F
SeqFold score									76.5	72.69	
PMF score		0.13	-0.06	0.23	0.1		0.42	_			-
Verify score		0.31	0	0.15	0.32		0.14	0.28			0.02
PSI- BLAST		3.20E-44	1.40E-42	6.40E-41	1.60E-44		9.00E-31	1.10E-28	1.10E-28	9.00E-28	9.00E-28
End AA		911	125	118	116		275	275	280	273	274
Start AA		19	19		61		216	216	216	217	217
Chain ID		1	1	L	L			∢	4	В	Ø
PDB ID		1fgv	11 gm	lvge	2fgw		Ibw 5	161	191	191	151
SEQ NO:		887	887	887	887		888	888	888	888	888

	Т								7
PDB annotation	TRANSCRIPTION 2 REGULATION	COMPLEX (GENE REGULATING PROTEIN/DNA) PAX, PRD, PAIRED DOMAIN, DNA-BINDING PROTEIN IPDN 14	COMPLEX (GENE REGULATING PROTEIN/DNA) PAX, PRD, PAIRED DOMAIN, DNA-BINDING PROTEIN IPDN 14	COMPLEX (GENE REGULATING PROTEIN/DNA) PAX, PRD, PAIRED DOMAIN, DNA-BINDING PROTEIN IPDN 14	DNA-BINDING DNA-BINDING, TRANSCRIPTION FACTOR, LFBI/ANF1, 2 HELIX-TURN-HELIX, DNA-BINDING DOMAIN	GENE REGULATION/DNA PAX, PAIRED DOMAIN, IRANSCRIPTION, PROTEIN-DNA INTERACTIONS, 2 GENE REGULATION/DNA	GENE REGULATION/DNA PAX, PARED DOMAIN, TRANSCRIPTION, PROTEIN-DNA INTERACTIONS, 2 GENE REGULATION/DNA	GENE REGULATION/DNA PAX, PAIRED DOMAIN, TRANSCRIPTION, PROTEIN-DNA INTERACTIONS, 2 GENE REGULATION/DNA	
	TRAD	COMPLE PROTEIN DOMAIN	COMPLE PROTEIN DOMAIN	COMPLE PROTEIN DOMAIN	TRAI LFBI DNA	PAIR PRO GEN	PROGEN	PAIR PRO REC	
Coumpound		PRD PAIRED DOMAIN; CHAIN: C; IPDN 4 DNA; CHAIN: A, B IPDN 5	PRD PAIRED DOMAIN; CHAIN: C; 1PDN 4 DNA; CHAIN: A, B 1PDN 5	PRD PAIRED DOMAIN; CHAIN: C; IPDN 4 DNA; CHAIN: A, B IPDN 5	LFBI/HNFI TRANSCRIPTION FACTOR; CHAIN: NULL;	HOMEOBOX PROTEIN PAX- 6. CHAIN: 4: 26 NUCLEOTIDE DNA; CHAIN: B; 26 NUCLEOTIDE DNA; CHAIN: C;	HOMEOBOX PROTEIN PAX- 6; CHAIN: 4; 26 NUCLEOTIDE DNA; CHAIN: B; 26 NUCLEOTIDE DNA; CHAIN: C;	HOMEOBOX PROTEIN PAX- 6; CHAIN: A; 26 NUCLEOTIDE DNA; CHAIN: B; 26 NUCLEOTIDE DNA; CHAIN: C;	
SeqFold score			168.62			143.36			
PMF		-		_	0.33				
Verify		0.57		99.0	0.31		0.61	0.48	
PSI- BLAST		9.60E-29	1.10E-63	1.10E-63	1.40E-30	1.60E-67	1.60E-31	1.60E-67	
End		152	160	160	280	691	152	169	
Start AA		35	35	35	209	34	35	36	
Chain		U	ပ	O		4	<	∢	
PDB		1pdn	1pdn	Ipdn	2lfb	брах	брах	ębax (
SEQ	į	888	888	888	888	888	888	888	

TABLE 6

TABLE 6 SEQ ID NO:	Position of Last Amino Acid of	Maximum Score	Mean Score
	Signal Peptide	0.002	0.021
445	21	0.993 0.975	0.931 0.962
446	14	0.986	0.606
447	42	0.908	0.703
448	18	0.967	0.778
449	27	0.992	0.946
450	30	0.997	0.973
452	17	0.907	0.575
454	32	0.931	0.672
455	27	0.988	0.755
456	40	0.986	0.916
457	26	0.920	0.750
458	18	0.946	0.790
459	15	0.993	0.931
460	21	0.942	0.644
461	47	0.886	0.712
463	24	0.985	0.865
464	36	0.965	0.679
465	42	0.980	0.946
466	43	0.969	0.858
467	27	0.959	0.793
469	26	0.983	0.687
470	45	0.981	0.821
471	25	0.998	0.963
472	30	0.977	0.915
473	18	0.949	0.644
474	27	0.949	0.768
475	23	0.947	0.901
476	19	0.936	0.628
477	15	0.956	0.893
478	17	0.942	0.720
479	17	0.952	0.730
480	17	0.970	0.916
481		0.975	0.962
483	14 47	0.955	0.727
486		0.991	0.952
488	23 42	0.986	0.606
495		0.971	0.594
496	11 29	0.896	0.743
502		0.908	0.703
509	18	0.959	0.908
510	20	0.957	0.858
512	24	0.967	0.778
516	35	0.991	0.851
517		0.939	0.722
518	26 47	0.983	0.640
519	30	0.992	0.946
522		0.974	0.924
538	16	0.997	0.973
550	42	0.947	0.588
551	30	0.981	0.684
555	30 32	0.907	0.575
576		0.973	0.927
577	26		0.672
578 589 590	27 40 38	0.931 0.988 0.985	0.672 0.755 0.775

			<u> </u>
595	20	0.938	0.818
611	18	0.920	0.750
615	25	0.949	0.775
616	33	0.995	0.835
617	15	0.946	0.790
623	19	0.921	0.819
627	21	0.993	0.931
634	20	0.961	0.674
635	28	0.954	0.648
645	47	0.942	0.644
647	31	0.962	0.776
650	16	0.949	0.782
651	14	0.963	0.613
654	20	0.984	0.958
670	24	0.886	0.712
673	17	0.934	0.753
678	36	0.985	0.865
695	23	0.954	0.754
707	42	0.965	0.679
708	22	0.979	0.667
709	24	0.984	0.851
710	17	0.911	0.745
717	25	0.980	0.946
718	35	0.988	0.871
726	27	0.969	0.858
730	17	0.981	0.844
741	22	0.937	0.871
755	17	0.890	0.668
764	26	0.950	0.793
768	. 32	0.958	0.827
771	45	0.983	0.687
773	39	0.997	0.801
776	17	0.945	0.650
787	32	0.983	0.835
789	25	0.981	0.821
792	31	0.966	0.815
796	22	0.887	0.572
797	19	0.941	0.691
807	30	0.998	0.963
808	18	0.977	0.915
809	18	0.977	0.915
811	27	0.959	0.827
812	16	0.925	0.734
815	19	0.934	0.564
816	21	0.960	0.858
818	27	0.949	0.644
821	27	0.943	0.758
823	27	0.908	0.728
833	23	0.913	0.768
837	19	0.947	0.901
841	22	0.967	0.826
845	15	0.936	0.628
846	20	0.975	0.840
851	31	0.985	0.908
852	19	0.965	0.922
853	39	0.984	0.743
857	17	0.956	0.893
858	21	0.957	0.868
861	22	0.975	0.866
868	21	0.942	0.736

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871	43	0.973	0.560
873	19	0.952	0.730
874	33	0.923	0.717
879	23	0.978	0.911
881	16	0.947	0.884
887	17	0.970	0.916

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TABLE 7

SEQ ID NO:	Position of Last Amino Acid of Signal Peptide	Maximum Score	Mean Score
445	21	0.993	0.931
446	14	0.975	0.962
447	42	0.986	0.606
448	18	0.908	0.703
449	24	0.967	0.778
450	30	0.992	0.946
452	17	0.997	0.973
454	32	0.907	0.575
455	27	0.931	0.672
456	40	0.988	0.755
457	26	0.986	0.916
458	18	0.920	0.750
459	15	0.946	0.790
460	21	0.993	0.931
461	47	0.942	0.644
463	24	0.886	0.712
464	36	0.985	0.865
	42	0.965	0.679
465	25	0.980	0.946
466	27	0.969	0.858
467	26	0.950	0.793
469		0.983	0.687
470	45	0.981	0.821
471	25	0.998	0.963
472	30 .		0.915
473	18	0.977	
474	27	0.949	0.644
475	23	0.913	0.768
476	19	0.947	0.901
477	15	0.936	0.628
478	17	0.956	0.893 ·
479	17	0.942	0.720
480	19	0.952	0.730
481	17	0.970	0.916
483	14	0.975	0.962
486	47	0.955	0.727
488	23	0.991	0.952
495	42	0.986	0.606
496	11	0.971	0.594
502	29	0.896	0.743
509	18	0.908	0.703
510	13	0.959	0.908
512	20	0.957	0.858
516	24	0.967	0.778
517	35	0.991	0.851
518	26	0.939	0.722
519	47	0.983	0.640
522	30	0.992	0.946
538	16	0.974	0.924
550	17	0.997	0.973
551	42	0.947	0.588
555	30	0.981	0.684
576	32	0.907	0.575
577	26	0.973	0.927
578	27	0.931	0.672
589	40	0.988	0.755

	I	0.005	0.775
590	38	0.985	0.775 0.818
595	20		0.750
611	. 18	0.920 0.949	0.775
615		0.995	0.835
616	33	0.946	0.790
617 623	15 19	0.940	0.790
		0.993	0.931
627	21	0.961	0.674
634	20	0.951	0.648
635	28	0.942	0.644
645	47	0.942	0.776
647	31		0.782
650	16	0.949	0.613
651	14		0.958
654	20	0.984	0.712
670	24		0.712
673	17	0.934	
678	36	0.985	0.865 0.754
695	23	0.954	
707	42	0.965 0.979	0.679 0.667
708 709	24	0.979	0.851
709	17	0.911	0.745
717	25	0.980	0.946
718	35	0.988	0.871
726	27	0.969	0.858
730	17	0.981	0.844
741	22	0.937	0.871
755	17	0.890	0.668
764	26	0.950	0.793
768	32	0.958	0.827
771	45	0.983	0.687
773	39	0.997	0.801
776	17	0.945	0.650
787	32	0.983	0.835
789	25	0.981	0.821
792	31	0.966	0.815
796	22	0.887	0.572
797	19	0.941	0.691
807	30	0.998	0.963
808	18	0.977	0.915
809	18	0.977	0.915
811	27	0.959	0.827
812	16	0.925	0.734
815	19	0.934	0.564
816	21	0.960	0.858
818	27	0.949	0.644
821	27	0.943	0.758
823	27	0.908	0.728
833	23	0.913	0.768
837	19	0.947	0.901
841	22	0.967	0.826
845	15	0.936	0.628
846	20	0.975	0.840
851	31	0.985	0.908
852	19	0.965	0.922
853	39	0.984	0.743
857	17	0.956	0.893
858	21	0.957	0.868
. 861	22	0.975	0.866

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868	21	0.942	0.736
871	43	0.973	0.560
873	19	0.952	0.730
874	33	0.923	0.717
879	23	0.978	0.911
881	16	0.947	0.884
887	17	0.970	0.916

TABLE 8

SEQ ID NO: of Nucleotide Sequence	SEQ ID NO: of Polypeptide Sequence	SEQ ID NO: in USSN 09/659,671
		·
1	445	2
2	446	. 5
3	447	6
4	448	7
5	449	8
6	450	9
7	451	11
8	452	12
9	453	13
10	454	14
11	455	15
12	456	16
13	457	17
14	458	18
15	459	19
16	460	20
17	461	21
18	462	23
19	463	24
20	464	25
21	465	28
22	466	29
23	467	30
24	468	31
25	469	34
26	470	35
27	471	37
28	472	38
29	473	39
	474	40
30	475	41
31	476	42
32	477	43
33		44
34	478	45
35	479	46
36	480	47
37	481	47
38	482	50
39	483	51
40	484	52
41	485	
42	486	53
43	487_	54
44	488	55
45	489	56
46	490	57
47	491	58
48	492	59
49	493	60
50	494 -	61
51	495	62
52	496	63
53	497	64
54	498	65

	100	
55	499 500	66
56		68
57	501	69
58	502	70
59	503	71
60	504	
61	505	72
62	506	73
63	507	74
64	508	75
65	509	76 .
66	510	77
67	511	78
68	512	79
69	513	80
70	514	81
71	515	82
72	516	83
73	517	85
. 74	518	86
75	519	88
76	520	89
77	521	90
78	522	91
79	523	92
80	524	93
81	525	94
82	526	95
	527	96
83 84	528	97
	529	98
85	530	99
86	531	100
87		101
88	532	102
89	533	105
90	534	
91	535	106
92	536	107
93	537	108
94	538	109
95	539	110
96	540	111
97	541	112
98	542	113
99	543	114
100	544	115
101	545	116
102	546	117
103	547	118
104	548	119
105	549	120
106	550	121
107	551	122
108	552	123
109	553	124
110	554	125
	555	126
111	556	127
112		127
113 114	557	128
11/1	558	129

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115	559	130	
116	560	131	
117	561	132	
118	562	133	
119	563	134	
120	564	135	
121	565	136	
122	566	137	
123	567	138	
124	568	139	
125	569	140	
126	570	141	
127	571	142	
128	572	143	
129	573	144	
130	574	145	
131	575	146 147	
132	576 577	147	
133 134	578	150	
134	579	151	
136	580	152	
137	581	153	
138	582	154	
139	583	155	
140	584	156	
141	585	157	
142	586	158	
143	587	159	
144	588	160	
145	589	161	
146	590	162	
147	591	163	
148	592	165	
149	593	166	
150	594	167	
151	595	168	
152	596 .	169	
153	597	. 170	
154	598	171	
155	599	172	
156	600	173	
157	601	174	
158	602	175 176	
159 160	604	177	
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161	605	· 178	
163	607	180	
164	608	181	
165	609	182	
166	610	183	
167	611	184	
168	612	185	
169	613	186	
170	614	187	
171	615	188	
172	616	189	
173	617	190	
174	618	191	
			

175	619	192
176	620	193
177	621	194
178	622	195
179	623	197
180	624	198
181	625	199
182	626	200
183	627	201
184	628	202
185	629	203
186	630	204
187	631	205
188	632	206
189	633	207
190	634	208
191	635	209
192	636	210
193	637	211
194	638	212
195	639	213
196	640	214
197	641	215
198	642	216
199	643	217
200	644	218
201	645	219
202	646	220
	647	221
203 204	648	222
	649	223
205	650	225
206	651	226
207		227
208	652	228
209	653	229
210	654	230
211	655	
212	656	231
213	657	232
214	658	233
215	659	234
216	660	235
217	661	236
218	662	237
219	663	238
220	664	239
221	665	240
222	666	241
223	667	242
224	668	243
225	669	244
226	670	245
227	671	246
228	. 672	247
229	673	248
230	674	249
231	675	250
232	676	251
233	677	252
234	678	253
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235	679	254
236	680	255
237	681	256
238	682	258
239	683	259
240	684	260
241	685	261
242	686	262
243	687	263
244	688	264
245	689	265
246	690	266
247	691	267
248	692	268
249	693	269
250	694	270
	695	271
251		272
252	696	273
253	697	
254	698	274
255	699	275
256	700	277
257	701	278
258	702	279
259	703	280
260	704	282
261	705	283
262	706	284
263	707	285
264	708	286
265	709	287
266	710	288
267	711	289
268	712	290
269	713	291
270	714	292
271	715	293
272	716	294
273	717	295
274	718	296
275	719	297
276	720	298
277	721	299
278	722	300
279	723	301
280	724	302
281	725	304
282	726	305
283	727	306
284	728	307
285	729	308
286	730	309
287	731	310
288	732	311
289	733	312
290	733	313
290	735	314
292	736	316
		317
293	737	
294	738	318

302		
295	739	319
296	740	320
297	741	321
298	742	322
299	743	323
300	744	324
301	745	325
302	746	326
303	747	327
304	748	328
305	749	329
306	750	330
307	751	331
308	752	332
309	753	333
310	754	334
311	755	335
312	756	336
313	757	337
314	758	338
315	759	339
316	760	340
317	761	341
318	762	342
319	763	343 344
320	764 765	345
321 322	766	346
323	767	347
324	768	348
325	769	349
326	770	350
327	771	351
328	772	352
329	773	353
330	774	354
331	775	355
332	776	356
333	777	357
334	778	358
335	779	359
336	780	360
337	781	361
338	782	362
339	783	363
340	784	364
341	785	366
342	786	367
343	787	368
344	788	369
345	789	370
346	790 701	371 372
347	791	373
348	792 793	374
349 350	794	376
	794	377
351 352	796	378
353	797	379
353	798	, 380
334	130	, 300

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355	799	381
356	800	382
357	801	383
358	802	384
359	803	385
360	804	386
361	805	387
362	806	388
363	807	389
364	808	390
365	809	191
366	810	392
367	811	393
368	812	395
369	813	396
370	814	397
371	815	398
372	816	399
373	817	400
374	818	401
375	819	402
376	820	403
377	821	404
378	822	406
379	823	407
380	824	408 .
381	825	409
382	826	410
383	827	411
384	828	412
385	829	413
386	830	414
387	831	416
388	832	417
389	833	418
390	834	419
391	835	420
392	836	421
393	837	422
	838	423
394	839	424
395	840	425
396		426
397	841	427
398	842	427
399	843	428
400	844	
401	845	430
402	846	431
403	847	432
404	848	433
405	849	435
406	850	436
407	851	437
408	852	438
409	853	439
410	854	440
411	855	441
412	856	442
413	857	443
414	858	445

415	859	446
416	860	447
417	861	448
418	862	449
419	863	450
420	864	451
421	865	452
422	866	453
423	867	454
424	868	455
425	869	456
426	870	457
427	871	458
428	872	459
429	873	460
430	874	461
431	875	462
432	876	463
433	877	464
434	878	465
435	879	466
436	880	467
437	881	468
438	882	469
439	883	470
440	884	471
441	885	472
442	886	473
443	887	474
444	888	475

WHAT IS CLAIMED IS:

- 1. An isolated polynucleotide comprising a nucleotide sequence selected from the group consisting of SEQ ID NO: 1-444, a mature protein coding portion of SEQ ID NO: 1-444, an active domain coding portion of SEQ ID NO: 1-444, and complementary sequences thereof.
- 2. An isolated polynucleotide encoding a polypeptide with biological activity, wherein said polynucleotide hybridizes to the polynucleotide of claim 1 under stringent hybridization conditions.
- 3. An isolated polynucleotide encoding a polypeptide with biological activity, wherein said polynucleotide has greater than about 90% sequence identity with the polynucleotide of claim 1.
- 4. The polynucleotide of claim 1 wherein said polynucleotide is DNA.
- 5. An isolated polynucleotide of claim 1 wherein said polynucleotide comprises the complementary sequences.
- 6. A vector comprising the polynucleotide of claim 1.
- 7. An expression vector comprising the polynucleotide of claim 1.
- 8. A host cell genetically engineered to comprise the polynucleotide of claim 1.
- 9. A host cell genetically engineered to comprise the polynucleotide of claim 1 operatively associated with a regulatory sequence that modulates expression of the polynucleotide in the host cell.
- 10. An isolated polypeptide, wherein the polypeptide is selected from the group consisting of:
 - (a) a polypeptide encoded by any one of the polynucleotides of claim 1;
 - (b) a polypeptide encoded by a polynucleotide hybridizing under stringent conditions with any one of SEQ ID NO: 1-444.

- 11. A composition comprising the polypeptide of claim 10 and a carrier.
- 12. An antibody directed against the polypeptide of claim 10.
- 13. A method for detecting the polynucleotide of claim 1 in a sample, comprising:
- a) contacting the sample with a compound that binds to and forms a complex with the polynucleotide of claim 1 for a period sufficient to form the complex; and
- b) detecting the complex, so that if a complex is detected, the polynucleotide of claim 1 is detected.
- 14. A method for detecting the polynucleotide of claim 1 in a sample, comprising:
- a) contacting the sample under stringent hybridization conditions with nucleic acid primers that anneal to the polynucleotide of claim 1 under such conditions;
- b) amplifying a product comprising at least a portion of the polynucleotide of claim 1; and
- c) detecting said product and thereby the polynucleotide of claim 1 in the sample.
- 15. The method of claim 14, wherein the polynucleotide is an RNA molecule and the method further comprises reverse transcribing an annealed RNA molecule into a cDNA polynucleotide.
- 16. A method for detecting the polypeptide of claim 10 in a sample, comprising:
- a) contacting the sample with a compound that binds to and forms a complex with the polypeptide under conditions and for a period sufficient to form the complex; and
- b) detecting formation of the complex, so that if a complex formation is detected, the polypeptide of claim 10 is detected.
- 17. A method for identifying a compound that binds to the polypeptide of claim 10, comprising:
- a) contacting the compound with the polypeptide of claim 10 under conditions sufficient to form a polypeptide/compound complex; and
- b) detecting the complex, so that if the polypeptide/compound complex is detected, a compound that binds to the polypeptide of claim 10 is identified.

- 18. A method for identifying a compound that binds to the polypeptide of claim 10, comprising:
- a) contacting the compound with the polypeptide of claim 10, in a cell, under conditions sufficient to form a polypeptide/compound complex, wherein the complex drives expression of a reporter gene sequence in the cell; and
- b) detecting the complex by detecting reporter gene sequence expression, so that if the polypeptide/compound complex is detected, a compound that binds to the polypeptide of claim 10 is identified.
- 19. A method of producing the polypeptide of claim 10, comprising,
- a) culturing a host cell comprising a polynucleotide sequence selected from SEQ ID NO: 1-444, a mature protein coding portion of SEQ ID NO: 1-444, an active domain coding portion of SEQ ID NO: 1-444, complementary sequences thereof and a polynucleotide sequence hybridizing under stringent conditions to SEQ ID NO: 1-444, under conditions sufficient to express the polypeptide in said cell; and
 - b) isolating the polypeptide from the cell culture or cells of step (a).
- 20. An isolated polypeptide comprising an amino acid sequence selected from the group consisting of any one of the polypeptides SEQ ID NO: 217-432, or 649-864, the mature protein portion thereof, or the active domain thereof.
- 21. The polypeptide of claim 20 wherein the polypeptide is provided on a polypeptide array.
- 22. A collection of polynucleotides, wherein the collection comprising the sequence information of at least one of SEQ ID NO: 1-444.
- The collection of claim 22, wherein the collection is provided on a nucleic acid array.
- 24. The collection of claim 23, wherein the array detects full-matches to any one of the polynucleotides in the collection.
- 25. The collection of claim 23, wherein the array detects mismatches to any one of the polynucleotides in the collection.

- 26. The collection of claim 22, wherein the collection is provided in a computer-readable format.
- 27. A method of treatment comprising administering to a mammalian subject in need thereof a therapeutic amount of a composition comprising a polypeptide of claim 10 or 20 and a pharmaceutically acceptable carrier.
- 28. A method of treatment comprising administering to a mammalian subject in need thereof a therapeutic amount of a composition comprising an antibody that specifically binds to a polypeptide of claim 10 or 20 and a pharmaceutically acceptable carrier.